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AFAPL-TR-67-114  
Part III, Volume II

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VAPORIZING AND ENDOTHERMIC FUELS  
FOR ADVANCED ENGINE APPLICATION

Part III. Studies of Thermal and Catalytic Reactions,  
Thermal Stabilities, and Combustion Properties  
of Hydrocarbon Fuels

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Shell Development Company,  
A Division of Shell Oil Company

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Calculation Procedure for Mach 8 Engine

## Station 1.

Use the following formulas from Ref. 12 to calculate the inlet area:

$$I_f = \frac{n_p (H_f) J}{V_1}$$

$$W_f = \frac{n_n (F_g)}{I_f}$$

$$A_1 = \frac{W_f}{(F/A)(\rho_1)(V_1)}$$

Assume:  $n_p = .412$

$n_n = .95$

$H_f = 18894$  Btu/lbm fuel

At 100,000 ft and  $M = 8$

$$T_1 = 420.1^\circ R$$

$$P_1 = 22.32 \text{ lb/ft}^2$$

$$\rho_1 = .000996 \text{ lbm/ft}^3$$

$$V_1 = 8050.96 \text{ ft/sec}$$

$$I_f = \frac{(.412)(18894)(778)}{(8051)}$$

$$I_f = 752.23 \text{ sec}$$

Assume:  $L/D = 6$

$$L = 450,000 \text{ lb}$$

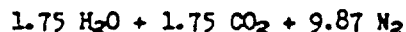
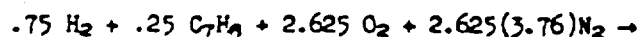
$$D = F_g = 75000 \text{ lb}$$

Therefore:

$$W_f = \frac{(.95)(75000)}{752.23}$$

$$W_f = 94.72 \text{ lbm/sec fuel}$$

To Calculate the stoichiometric fuel-air ratio (F/A) assume the fuel is MCH converted to  $H_2$  and  $C_7H_8$ :



The average molecular weight of the fuel: 25.25

$$F/A = \frac{(1 \text{ mole of fuel})(25.25 \text{ lbm molecular weight})}{(9.87 + 2.625 \text{ moles air})(28.95 \text{ lbm molecular weight})}$$

$$F/A = .0698$$

Assuming E.R. = 1.0

$$A_1 = \frac{94.72}{(.0698)(.000996)(5051)}$$

$$A_1 = 169.23 \text{ ft}^2$$

Station 3.

Use the Dugger Equation for the total pressure drop from Station 1 to 3:

$$n_d = 1 - \frac{\left(\frac{P_{t1}}{P_{t3}}\right)^{\bar{\gamma}-1/\bar{\gamma}} - 1}{((\bar{\gamma} - 1)/2)M_1^2}$$

Assume:  $n_d = .959$

$$\bar{\gamma} = 1.4$$

then

$$\frac{\bar{\gamma} - 1}{\bar{\gamma}} = \frac{1.4 - 1}{1.4} = .286$$

$$\frac{(P_{t1}/P_{t3})^{.286} - 1}{((1.4 - 1)/2)(8)^2} = 1 - .959$$

$$(P_{t1}/P_{t3})^{.286} = 1.5248$$

$$(P_{t1}/P_{t3}) = 4.36$$

From the isentropic flow tables at  $M = 8$

$$P_1/P_{t1} = .000102$$

$$P_{t1} = \frac{22.32 \text{ lb/ft}^2}{(.000102)(144 \text{ in}^2/\text{ft}^2)}$$

$$P_{t1} = 1519 \text{ lb/in}^2$$

$$P_{t3} = \frac{1512 \text{ lb/in}^2}{4.30}$$

$$P_{t3} = 348 \text{ lb/in}^2$$

Assume:

$$M_3 = 2.5 \quad \text{then}$$

From the isentropic flow tables at  $M = 2.5$

$$P_3/P_{t3} = .05853$$

$$P_3 = 20.4 \text{ lb/in}^2$$

Assume the enthalpy at 1 and 3 are equal:

$$h_{t1} = h_{t3} = h_3 + \frac{(M_3 49.1 \sqrt{T_3})^2}{2g_c J}$$

$$h_{t1} = 100.32 + \frac{((8)(49.1)\sqrt{420.1})^2}{(2)(32.2)(778)}$$

$$h_{t1} = 1394 \text{ Btu/lbm}$$

By trial and error

$$T_3 = 2500^\circ\text{R}$$

$$h_{t3} = 732.33 + \frac{((2.5)49.1\sqrt{2500})^2}{(2)(32.2)(778)}$$

$$h_{t3} = 1395 \quad \text{close enough}$$

$$\rho_3 = \frac{(20.4)(144)}{(53.34)(2500)}$$

$$\rho_3 = .0220 \text{ lbm/ft}^3$$

Station 5.

By an iterative procedure calculate the heat transfer from the combustion area assuming  $2000^\circ\text{R}$  wall temperature and thus the total temperature rise. Then using a Rayleigh line relationship the remaining conditions at Station 5 may be determined.

First calculate the flow area and heat transfer area:

$$\dot{m} = \rho_3 A_3 V_3 = \rho_1 A_1 V_1$$

$$V_3 = (2.5)(49.1)\sqrt{2500}$$

$$V_3 = 6140$$

$$A_3 = \frac{\rho_1 A_1 V_1}{\rho_3 V_3} = \frac{(0.000006)(169)(8051)}{(0.0220)(6140)}$$

$$A_3 = 10.0$$

$$10.0 = \pi r_1^2 - \pi r_3^2 = 169 - \pi r_3^2$$

$$r_3 = 7.11 \text{ ft}$$

Hydraulic Diameter  $D_h$

$$D_h = \frac{4A_3}{P_w} = \frac{4(10.0)}{2\pi(7.11) + 2\pi(7.28)}$$

$$D_h = .444$$

Assume time for combustion and mixing can be accomplished in  $2 \mu\text{sec} = .002 \text{ sec}$  and the average velocity in the combustor is 5000 ft/sec. The length of the combustor then is L

$$L = (.002)(5000)$$

$$L = 10 \text{ ft}$$

Forming a heat balance on the combustion area

$$(\dot{h}_p' - \dot{h}_{p0}') - (\dot{h}_R' - \dot{h}_{R0}') - \dot{q} = -\dot{h}_{RP0}$$

where:  $\dot{h}_R'$  = enthalpy of the reactants

$\dot{h}_p'$  = enthalpy of the products

$\dot{h}_{RP0}$  = heat of combustion

$\dot{q}$  = heat transferred per lbm fuel

Heat of combustion calculation:

$$\dot{h}_{RP0} = \dot{m}_{C_7H_8} (\dot{h}_{RP0})_{C_7H_8} + \dot{m}_{H_2} (\dot{h}_{RP0})_{H_2}$$

$$\dot{m}_{C_7H_8} = .938 \text{ lbm } C_7H_8/\text{lbm fuel}$$

$$\dot{m}_{H_2} = .062 \text{ lbm } H_2/\text{lbm fuel}$$

$$(h_{RfO})_{C_7H_8} = -17601 \text{ Btu/lbm}$$

$$(h_{RfO})_{C_7H_8} = -51593 \text{ Btu/lbm}$$

$$h_{RfO} = (.938)(-17601) + (.062)(-51593)$$

$$h_{RfO} = -19702.5 \text{ Btu/lbm fuel}$$

The reactants:

$$h_R' - h_{RfO}' = (h' - h_o')_{air} + (h' - h_o')_{fuel}$$

$$(h' - h_o')_{air} = \frac{M_{air}}{M_{fuel}} (h_{2500} - h_{537})$$

$$= \frac{(2.625 + 9.87) 28.92}{25.25} (645.78 - 128.34)$$

$$(h' - h_o')_{air} = 7412.8 \text{ Btu/lbm fuel}$$

$$(h' - h_o')_{fuel} = C_{Pfuel}(t - t_o)$$

Assume:  $C_{Pfuel} = .766 \text{ Btu/lbm} \cdot ^\circ\text{F}$

$$t = 900 ^\circ\text{F}$$

$$(h' - h_o')_{fuel} = (.766)(900 - 77)$$

$$(h' - h_o')_{fuel} = 631 \text{ Btu/lbm fuel}$$

$$h_R' - h_{RfO}' = 8043.8 \text{ Btu/lbm fuel}$$

The products:

Calculate enthalpy of products as function of temperature and make plot.

Constituent	n	$h_{537}$	$h_{4000}$	$n\Delta h$	$h_{5000}$	$n\Delta h$	$h_{5360}$	$n\Delta h$
CO <sub>2</sub>	1.75	4030	49231	79102	54000	104948	69677	114881
H <sub>2</sub> O	1.75	4258	40489	63404	53327	85871	58339	94991
N <sub>2</sub>	9.87	3730	31329	272406	40080	358773	43436	391898
				414913		549591		601771

$$(h_P' - h_{PfO}')_t = \frac{n\Delta h}{M_t} = \frac{n\Delta h}{25.25}$$



$T_{asp},$ °R	$(h_p' - h_{p_c}')_c$ Btu/lbm Fuel	
4000	16432	} See plot on following page (Figure T2)
5000	21766	
5380	23833	

Estimate of Heat Transfer From the Combustor:

Procedure:

1. Assume Temperature at Station 5
2. Calculate Mach Number and pressure at Station 5 from Rayleigh line
3. Assume linear variation of pressure and temperature over length of combustor
4. Calculate density, velocity, viscosity, Reynolds Number, Nusselt Number, correction constant for high velocity heat transfer, thermal conductivity, heat transfer coefficient, adiabatic wall temperature and the final product,  $hP\Delta t$ , where  $P$  is the wetted perimeter,  $\Delta t = t_{aw} - t_w$  and  $t_w = 2000^\circ R$ , and  $h$  is the local heat transfer coefficient.
5. From curve of  $hP\Delta t$  vs  $x$  calculate heat transferred;

$$q = - \int_{x=0}^{x=L} hP\Delta t dx$$

6. From  $q$  and  $(h' - h_0')_p$ ,  $(h' - h_0')_R$  and  $h_{p_0}$  calculate temperature at Station 5.
7. Return to 2 until temperature used to calculate values and calculated temperature are equal.

Example:

Station 3:

$M = 2.5$

$P = 20.4$

$T = 2500^\circ R$

$(P/P_0) = .24616$

$(T/T_0) = .3787$

Assume  $T_5 = 5637.5^\circ R$

$$(T/T_0)_3 = (T/T_0)_2 T_2/T_3 = (.5787) \frac{(5637.5)}{(2500.)}$$

$$(T/T_0)_3 = .8540$$

From Rayleigh Line:

$$M_3 = 1.31$$

$$(P/P_0)_3 = .70535$$

$$P_3 = (P/P_0)_3 / (P/P_0)_2 P_2 = (.70535) / (.24616) (20.4)$$

$$P_3 = 58.45 \text{ lb/in}^2$$

x	P	T	$\rho$	V	$\mu$ $\times 10^{-5}$	Re $\times 10^6$	$Nu_1$ ( $M=0$ )	M	c	$Nu_1$	$\epsilon$
0	20.4	2500	.022	6150	3.4	1.76	1330	2.50	.65	864.5	.048
2	28.0	3110	.024	5633	3.8	1.58	1270	2.06	.75	952.5	.056
4	35.6	3740	.026	5200	4.2	1.43	1220	1.73	.81	988.2	.062
6	43.3	4370	.027	5007	4.6	1.30	1170	1.54	.84	982.8	.0685
8	51.0	5000	.028	4872	4.9	1.24	1145	1.40	.88	1007.6	.074
10	58.45	5637.5	.028	4828	5.2	1.17	1110	1.31	.89	987.9	.079

• Extrapolated (based on air at low pressures)

x	Pr	Pr 1/3	$T_{AW}/T$	$T_{AW}$	$\Delta t$	h	$hPr\Delta t$
0	.73	.90	2.125	5312.5	3312.5	93.4	$2.80 \times 10^7$
2	.75	.91	1.77	5504.7	3504.7	120.1	$3.81 \times 10^7$
4	.79	.92	1.55	5797.0	3797.0	138.0	$4.74 \times 10^7$
6	.90	1.0	1.47	6423.9	4423.9	151.6	$6.07 \times 10^7$
8	1.00	1.0	1.39	6950.0	4950.0	167.9	$7.52 \times 10^7$
10	1.00	1.0	1.34	7554.3	5554.3	175.8	$8.84 \times 10^7$

In the above calculations the following expressions were used:

$$\rho = P/RT \quad \text{where } R = 53.34$$

$$V = \rho_1/V_1/\rho = 135.2/\rho$$

$$Re = \frac{V D_n \rho}{\mu}$$

$$Nu_1 = f(Re, Pr) \quad \text{from Kays}^{14)} \text{ expression for turbulent flow inside concentric annuli at } r_i/r_o = 1.0 \text{ (See Figure 73)}$$

$$M = V/(49.1\sqrt{T})$$

$$C = \text{From Kays page 13.28 for affect of high velocity on Stanton No.}$$

$$M_1 = C M_{u1}$$

$$h = \frac{M_{1k}}{D_n}$$

$$T_{AW}/T = 1 + \frac{Pr^{1/3}}{2} (\gamma - 1) M^2 \quad \text{where } \gamma = 1.4$$

$$\Delta t = (T_{AW} - T_W) = (T_{AW} - 2000)$$

$$P = 2\pi r_1 + 2\pi r_0 = 90.5 \text{ ft}$$

Then:  $q' = - \int_{x=0}^{x=L} (h P \Delta t)_x dx$  This expression is integrated graphically from Figure 7b

$$q' = - 5.575 \times 10^8 \text{ Btu/hr}$$

$$\dot{m}_{air} = \rho_1 A_1 V_1 = 4.879 \times 10^8 \text{ lbm/hr}$$

$$q = - \frac{5.575 \times 10^8}{4.879 \times 10^8}$$

$$q = - 1.1426 \times 10^2 \text{ Btu/lbm air}$$

$$q' = - \frac{1.1426 \times 10^2}{.0698}$$

$$q' = - 1637 \text{ Btu/lbm fuel}$$

Now to calculate the combustion temperature:

$$(h' - h_o')_p - (h' - h_o')_R - q' = - h_{RPo}$$

$$(h' - h_o')_p - 8043.8 + 1637 = - (-19708.5)$$

$$\therefore (h' - h_o)_p = 26115.3 \text{ Btu/lbm fuel}$$

From curve in Figure 7j this gives a temperature:

$$T_3 = 5800^\circ R \text{ which does not agree with assume temp, } \therefore \text{ recalculate}$$

However, by plotting the calculated temperatures vs the assumed temperatures we can find the point they are equal. In addition by plotting the heat transferred vs the assumed temperature we can obtain the actual heat transfer at the actual temperature - see curves in Figure 7j.

This yields:

$T_3 = 5791^\circ R$ $q' = 1727 \text{ Btu/lbm fuel}$
--

$$\text{Then } (T/T_0)_5 = (.3787) \frac{(5791)}{(2500)}$$

$$(T/T_0)_5 = .8774 \quad \text{from Rayleigh line we get}$$

$$M_5 = 1.265$$

$$(P/P_0)_5 = .740$$

$$P_5 = \frac{(.740)}{(.24616)} (20.4)$$

$$P_5 = 61.3 \text{ lb/in}^2$$

## Station 6.

- Assume: 1. Isentropic expansion from 5 to 6.  
2.  $A_6 = 313.58 \text{ ft}^2$  ( $D = 20 \text{ ft}$ )

$$(A/A_0)_6 = (A/A_0)_5 \frac{A_5}{A_6} = \frac{(1.052)(313.58)}{10.0}$$

$$(A/A_0)_6 = 32.89$$

$$M_6 = 5.28$$

$$(P/P_0)_6 = .001538$$

$$P_6 = \frac{(.001538)(61.3)}{.377}$$

$$P_6 = .250 \text{ lb/in}^2$$

$$T_6 = \frac{(T/T_0)_6}{(T/T_0)_5} T_5$$

$$T_6 = \frac{(.15415)(5791)}{.758}$$

$$T_6 = 1177.88^\circ \text{R}$$

$$V = (5.28)(49.1) \sqrt{1177.88}$$

$$V_6 = 8897.48 \text{ ft/sec}$$

$$p_e = \frac{36.05}{(53.34)(1177.88)}$$

$$\rho_a = .000574 \text{ lbm/ft}^3$$

Calculate Thrust, Specific Impulse, and Overall Efficiency:

$$\left. \begin{aligned} \text{Thrust} = F_g &= f_e - f_1 - P_1(A_e - A_1) \\ f_e &= A_e \left( P_e + n_n \frac{\rho_a V_e^2}{g_c} \right) \\ f_1 &= A_1 \left( P_1 + \frac{\rho_1 V_1^2}{g_c} \right) \\ n_n &= .95 \end{aligned} \right\} \text{From Tuggeris}$$

$$f_1 = 169 \left( 22.32 + \frac{(.000996)(8051)^2}{32.2} \right)$$

$$f_1 = 342644.1 \text{ lb}$$

$$f_e = 313.58 \left( 36.05 + \frac{.95(.000574)(8897.48)^2}{32.2} \right)$$

$$f_e = 431703.9 \text{ lb}$$

$$F_g = 431703.9 - 342644.1 - 22.32(313.58 - 169.)$$

$$F_g = 85833.22 \text{ lb}$$

Specific Impulse:

$$I_f = \frac{n_n(F_g)}{W_f}$$

$$W_f = f_{o1} V_1 A_1 = (.0698)(.000996)(169.)(8051)$$

$$W_f = 94.6 \text{ lbm/sec fuel}$$

$$I_f = \frac{(.95)(85833.22)}{94.6}$$

$$I_f = 861.96 \text{ sec}$$

Overall Efficiency:

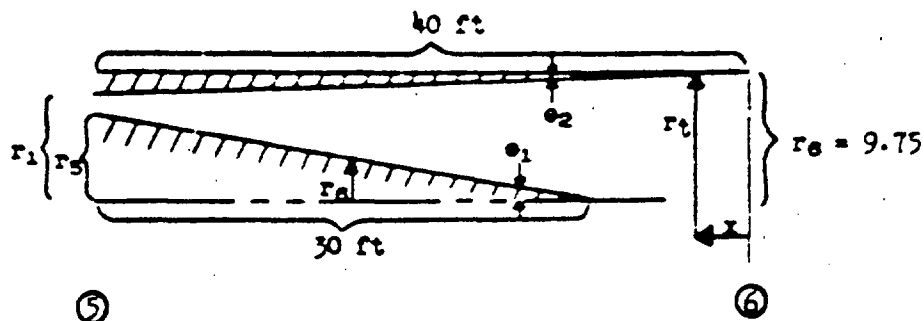
$$\eta_o = \frac{I_f V_1}{E_f + V_1^2/2g_c}$$

$$n_e = \frac{(861.96)(8050.96)}{(19708.5 + \frac{(8050.96)^2}{(2)(32.2)(778)}) 778}$$

$$n_e = .4247$$

### Estimate of Heat Transfer From Nozzle:

Consider the nozzle below: (Note: the values below are not the final but a first estimate)



Flow Area:

$$r_2 - r_1 = 9.75 - 7.28 = 2.47$$

$$\tan e_2 = \frac{2.47}{40} = .06175 \quad \tan e_1 = \frac{7.11}{30.0} = .237$$

$$e_2 = 3\frac{1}{2}^\circ$$

$$\text{From } x = 0 \text{ to } x = 10 \quad \text{where } r_t = r_2 - x \tan e_2 = 9.75 - x (.06175)$$

$$\text{From } x = 10 \text{ to } x = 40 \quad A_x = \underbrace{\pi r_t^2}_{A_t} - \underbrace{\pi r_b^2}_{A_b} \quad r_b = (x - 10) \tan e_1 = (x - 10) .237$$

$$x' = 40 - x$$

x'	x	r <sub>t</sub>	A <sub>t</sub>	r <sub>b</sub>	A <sub>b</sub>	A <sub>x</sub>
40	0	9.75	298.8	0	0	298.8
30	10	9.13	262.0	0	0	262.0
20	20	8.52	227.8	2.37	17.6	210.2
10	30	7.90	195.9	4.74	70.6	125.3
5	35	7.60	180.9	5.92	110.3	70.6
4	36	7.53	178.0	6.16	119.3	58.7
3	37	7.47	175.1	6.40	128.6	46.5
2	38	7.40	172.2	6.64	138.3	33.9
1	39	7.34	169.3	6.87	148.4	20.9

Assuming isentropic expansion to each point:

$$(A/A_0)_X = \frac{(A/A_0)_S A_X}{A_S} = \frac{1.1}{10.3} = .10967 A_X \quad (\text{ft}^2)$$

This yields a Mach Number from Isentropic Flow Tables.

$$T_X = \frac{(T/T_0)_X}{(T/T_0)_S} T_S = \frac{(T/T_0)_X 5465}{.726} = 7527.5 (T/T_0)_X \quad (^\circ\text{R})$$

$$P_X = \frac{(P/P_0)_X}{(P/P_0)_S} P_S = \frac{(P/P_0)_X 54}{.323} = 167 (P/P_0)_X \quad (\text{lb/in}^2)$$

$$V_X = M_X 49.1 \sqrt{T_X} \quad (\text{ft/sec})$$

$$\rho_X = \frac{P_X 144}{53.34 T_X} = 2.7 P_X / T_X \quad (\text{lbm/ft}^3)$$

x'	(A/A <sub>0</sub> ) <sub>X</sub>	M	(P/P <sub>0</sub> ) <sub>X</sub>	P <sub>X</sub>	(T/T <sub>0</sub> ) <sub>X</sub>	T <sub>X</sub>	V <sub>X</sub>	ρ <sub>X</sub>	μ
1	2.292	2.35	.074	12.38	.475	3525	6851	.00948	4.4 x 10 <sup>-5</sup>
2	3.718	2.86	.034	5.68	.379	2815	7451	.00544	3.93
3	5.10	3.20	.020	3.34	.328	2430	7745	.00371	3.68
4	6.44	3.45	.014	2.34	.290	2150	7855	.00294	3.47
5	7.74	3.64	.0108	1.81	.274	2063	8118	.00236	3.40
10	13.74	4.28	.0050	.836	.220	1656	8552	.00136	3.07
20	23.05	4.90	.00213	.356	.172	1294	8654	.00074	2.75
30	28.73	5.10	.00165	.276	.160	1200	8674	.00062	2.66
40	32.89	5.28	.00154	.257	.154	1160	8831	.00060	2.62

Assume the heat transfer relation is turbulent flow flat plate.

$$Nu_X = c(.332(Pr)^{1/3}(Re_X)^{1/2}) = cNu_1$$

$$Re_X = \frac{V_X \rho}{\mu}$$

c = correction constant for high speed flow (Kays, <sup>14</sup> p. 13.28)

$$h = \frac{Nu_X k}{X'} \quad (\text{Btu/hr-ft}^2\text{-}^\circ\text{F})$$

x'	Re	Re <sup>1/2</sup>	Pr	Pr <sup>1/3</sup>	Nu <sub>1</sub>	c	Nu <sub>X</sub>	k	h
1	1.475 x 10 <sup>6</sup>	1.214 x 10 <sup>3</sup>	.83	.94	378.9	.70	265	.065	17.2
2	2.06	1.435	.76	.91	433.5	.63	273	.058	7.9
3	2.34	1.530	.74	.90	457.2	.58	265	.054	4.8

(Continued)

$x'$	$R_e$	$R_e^{1/2}$	$Pr$	$Pr^{1/3}$	$Nu'$	$c$	$Nu_x$	$k$	$h$
4	2.66	1.63	.73	.90	487.0	.55	268	.050	3.4
5	2.82	1.68	.73	.90	502.0	.54	271	.049	2.7
10	3.79	1.946	.71	.89	575.0	.45	259	.044	1.1
20	4.66	2.16	.70	.89	638.2	.38	243	.038	.46
30	6.06	2.46	.70	.89	726.9	.37	269	.037	.33
40	8.06	2.84	.696	.89	839.2	.36	302	.036	.27

$$T_{ax} = T_x \left( 1 + \frac{r_c}{2} (\gamma - 1) M^2 \right)$$

$$\gamma = 1.4$$

$$r_c = Pr^{1/3}$$

$$\Delta t = T_{AW} - T_W = (T_{AW} - 2000^\circ R)$$

$$P = 2\pi r_t + 2\pi r_b$$

$x'$	$T_{AW}$	$\Delta t$	$P$	$hP\Delta t$
1	7184.8	5184.8	89.3	$7.96 \times 10^6$
2	6959.6	4959.6	88.2	3.46
3	6909.0	4909.0	87.1	2.05
4	6756.3	4756.3	86.0	1.39
5	6983.1	4983.1	84.9	1.14
10	7116.3	5116.3	79.4	.447
20	6886.4	4886.4	68.4	.154
30	6618.2	4618.2	57.4	.091
40	6981.0	4981.0	61.3	.082

Then:

$$q_N = \int_{x'=0}^{x'=40} (hP\Delta t)_x dx'$$

Integrating graphically from curve of  $x'$  vs  $hP\Delta t$  in Figure 76 we get,

$$q_N = 47.8 \times 10^6 \text{ Btu/hr}$$

$$q_N = \frac{47.8 \times 10^6 \text{ Btu/hr}}{4.879 \times 10^6 \frac{\text{lbm air}}{\text{hr}}}$$



$$q'_N = 9.797 \text{ Btu/lbm air}$$

$$q'_N = \frac{9.797}{.0098} \text{ lbm fuel/lbm air}$$

$$q'_N = 140.4 \text{ Btu/lbm fuel}$$

$$q'_T = q'_N + q'_c$$

$$q'_T = 1867.4 \text{ Btu/lbm fuel}$$

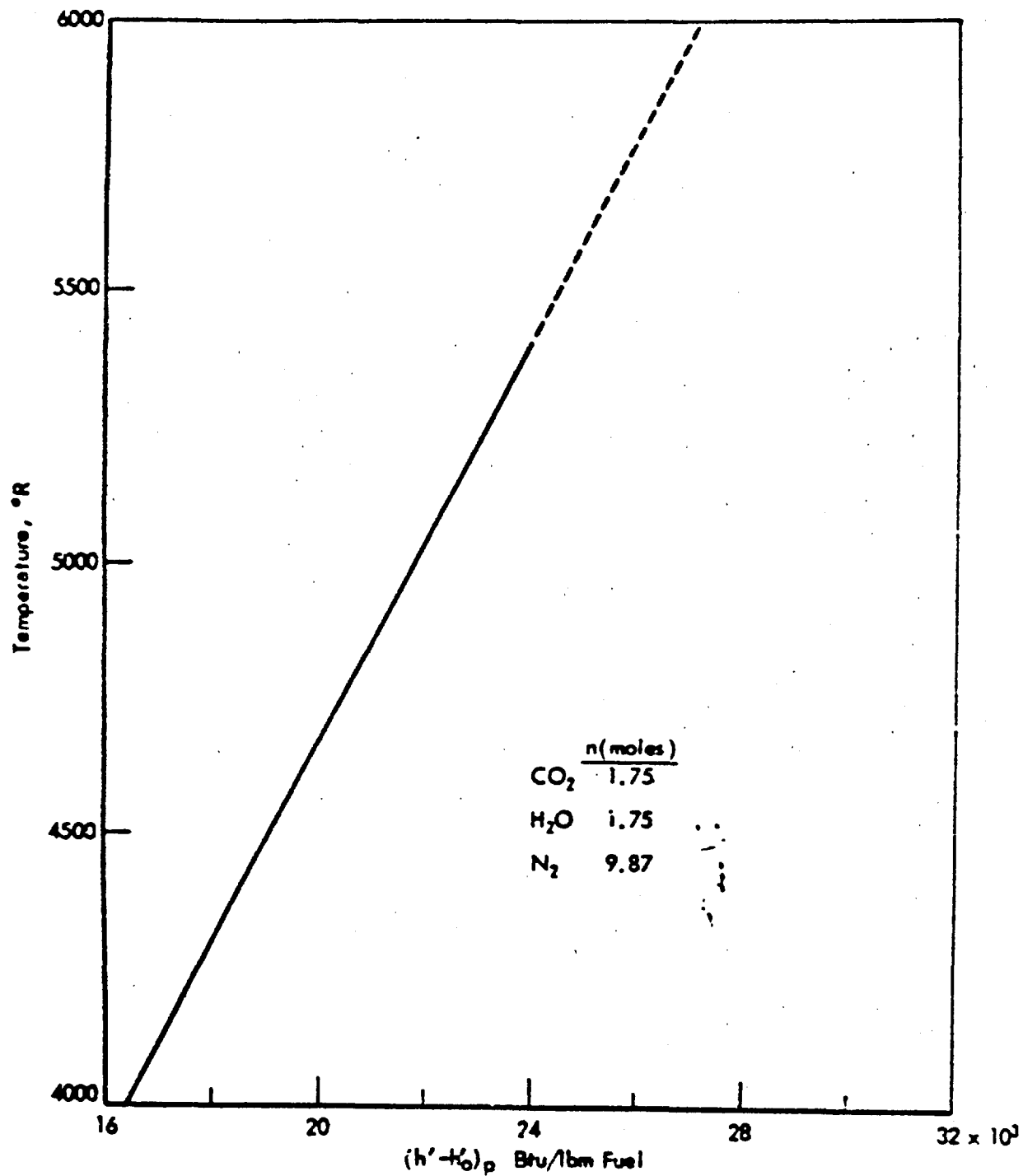


Figure 72. ENTHALPY OF PRODUCTS OF COMBUSTION

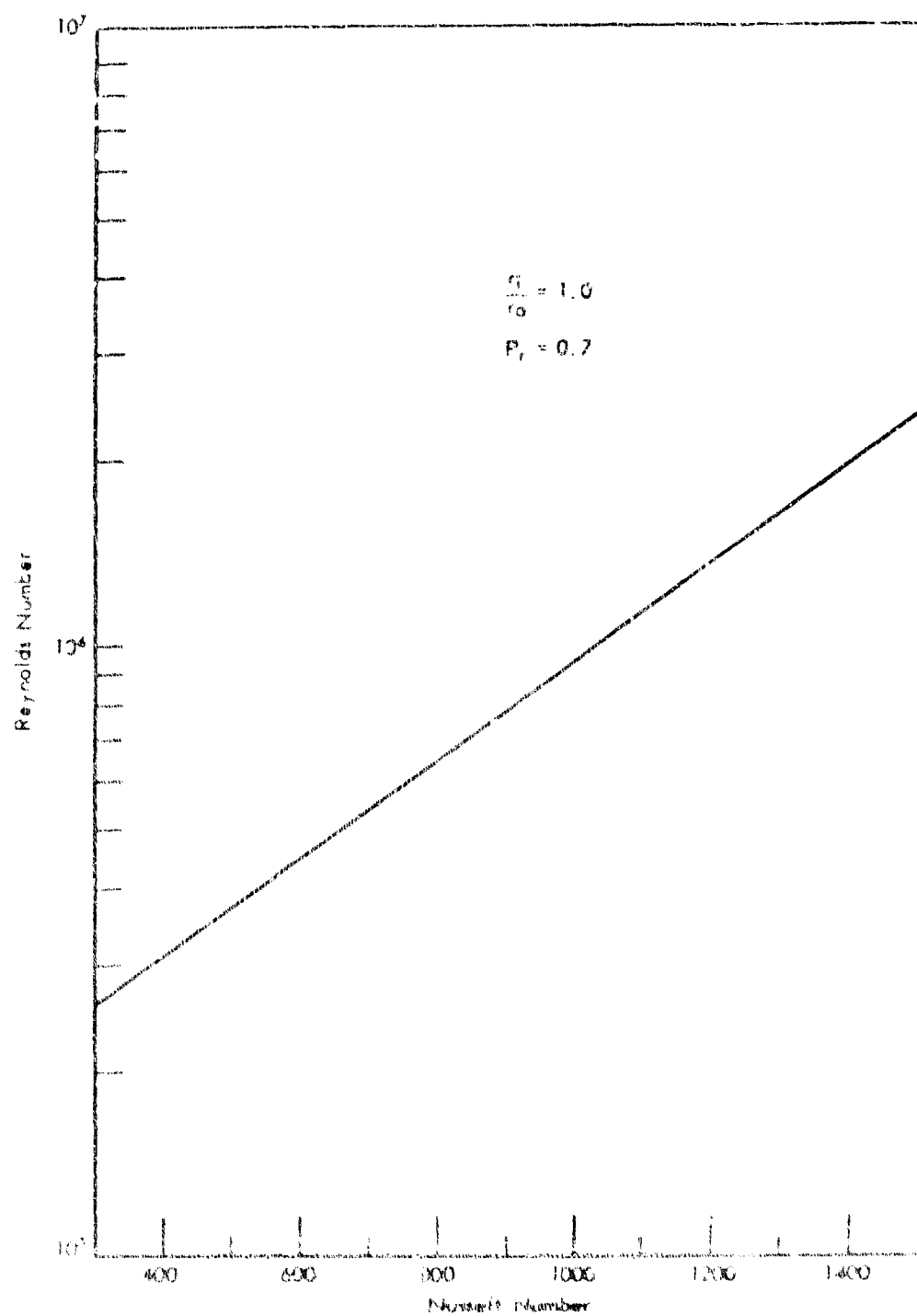


Figure 73. TURBULENT ANNULAR FLOW  
From Kay, Table 7-7

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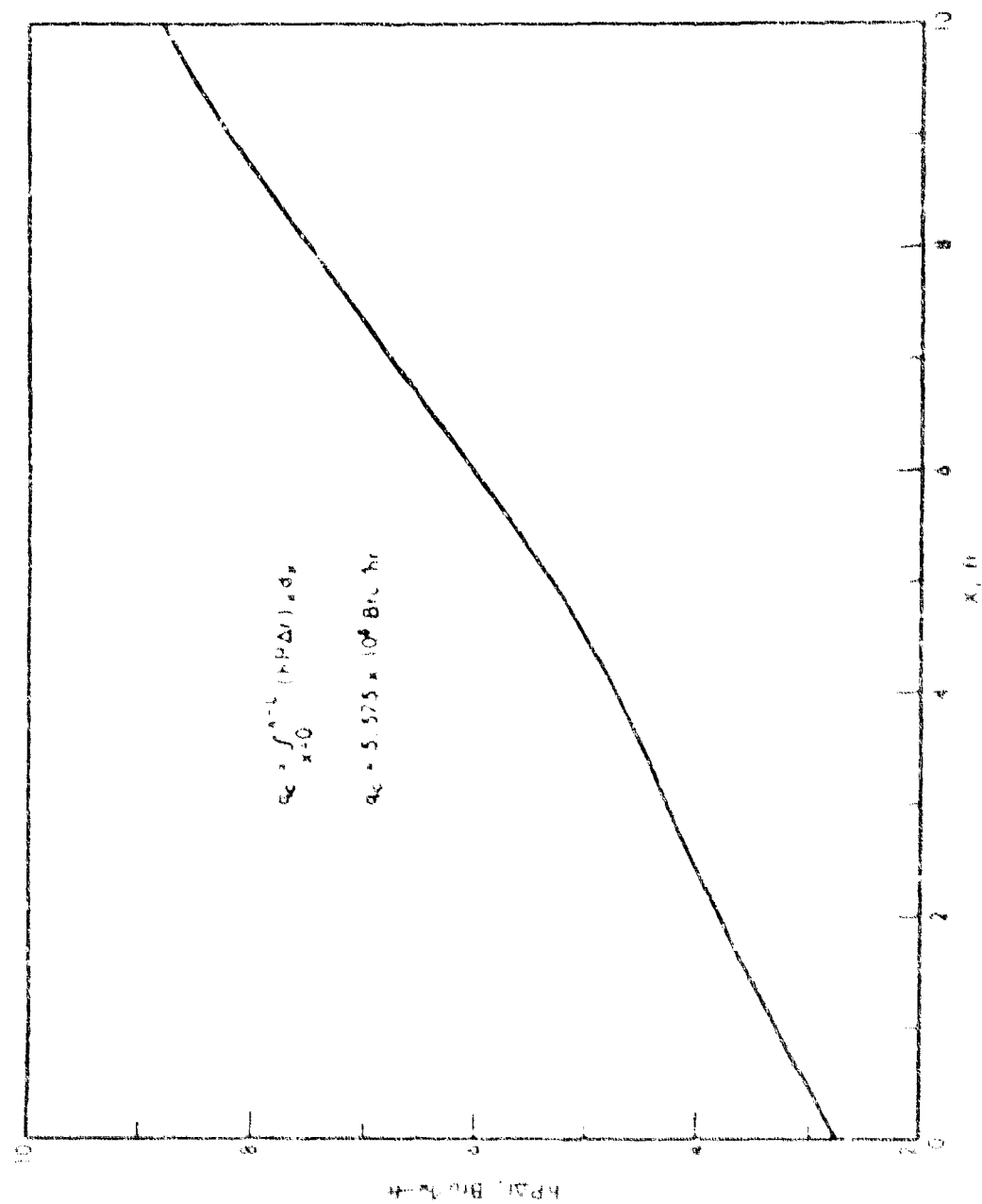


Figure 24. COMBUSTOR HEAT FLUX DISTRIBUTION

END

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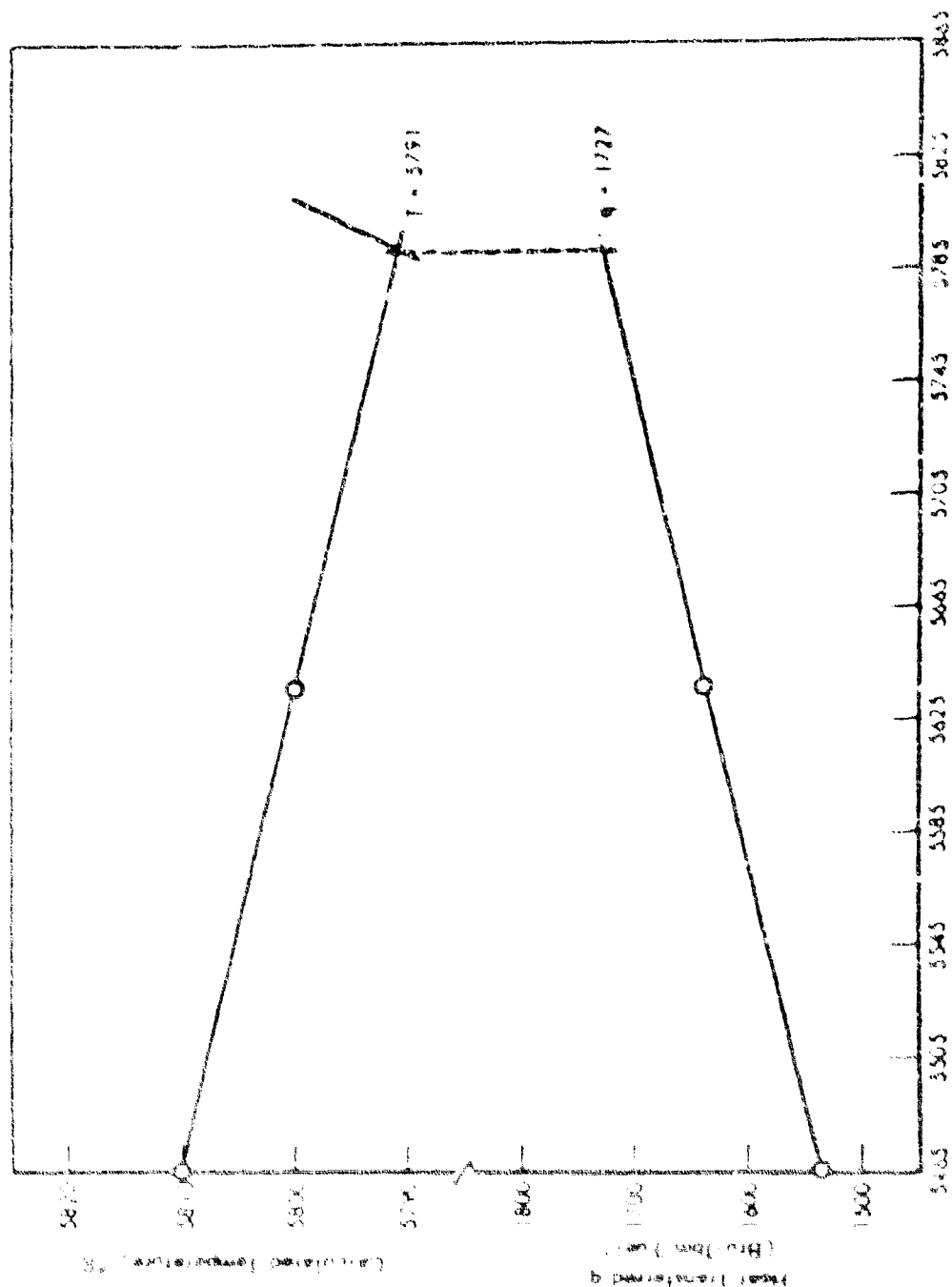


Figure 75. HEAT TRANSFER CONDITIONS

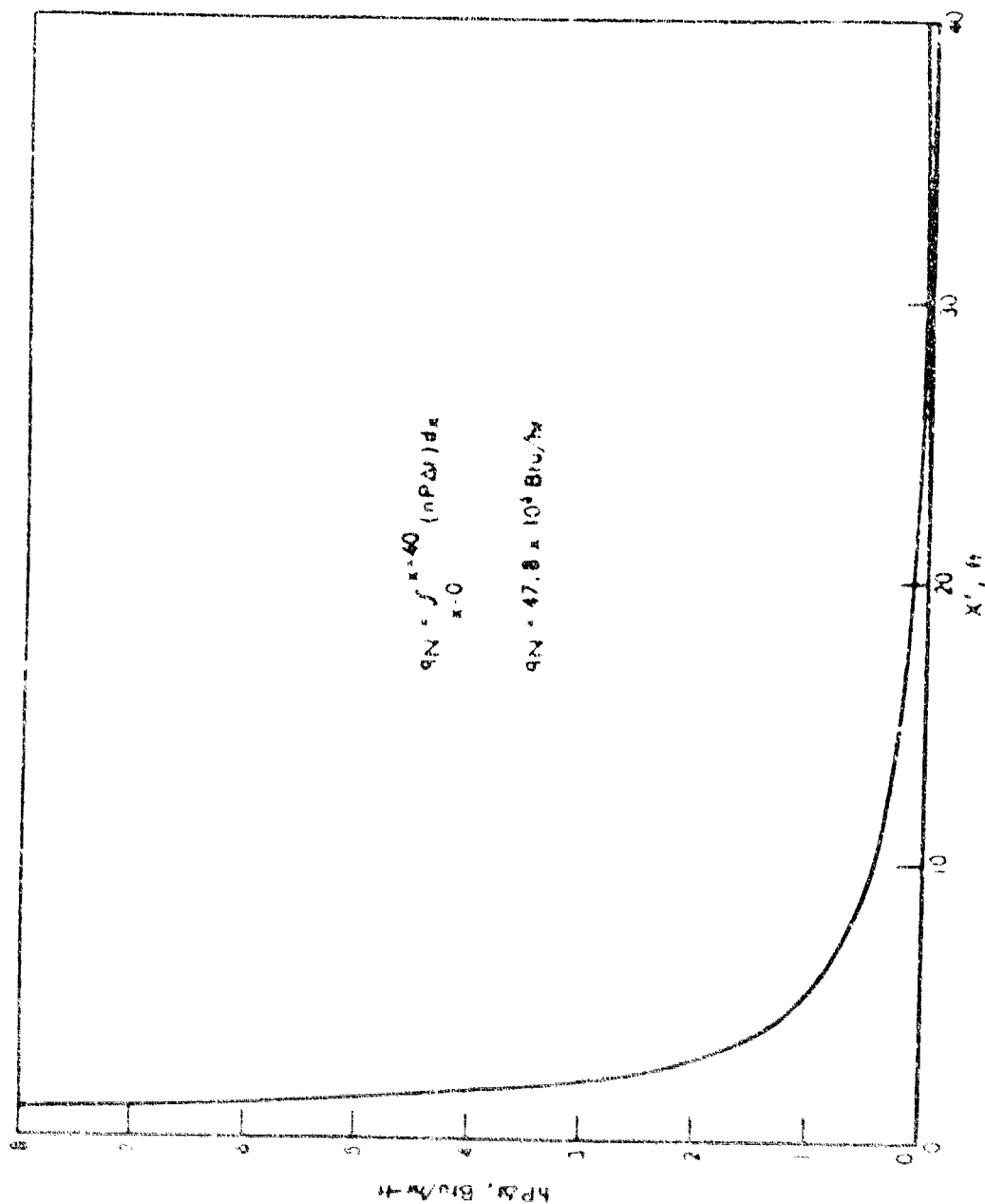


Figure 76. NOZZLE HEAT FLUX DISTRIBUTION

Table 112. DEHYDROGENATION OF MCH OVER AEROSOL-PHF-1

Pressure: 1 atm Catalyst Volume: 7 ml  
Block Temperature: 842 °F Reaction Time: 30 Minutes

Run Number 11018-	35	36-1	36-2	37
LHSV	5	15	30	50
Catalyst Bed Profile, °F	752 808-06 826-24 835-33	655-60 680-75 725-25 772-66	662-777 640-759 658-741 644-734	812-38 <sup>d)</sup> 820-36 813-35 808-35
Reactor Wall Temperature, °F	824-22	74-59	730-94	
CH <sub>4</sub> max	0	5	135	27 <sup>d)</sup>
Product Analysis, %				
Benzene	1.8	0.2	0.0	0.0
U <sub>1</sub> <sup>a)</sup>	0.0	0.1	0.4	0.6
MCH	1.0	3.7	41.6	96.1
U <sub>2</sub> <sup>b)</sup>	0.0	0.0	0.0	1.0
Toluene	97.1	96.0	58.0	1.2
U <sub>3</sub> <sup>c)</sup>	0.1	0.0	0.0	1.1
MCH Conversion, %	99.0	96.3	58.4	3.9

- a) Unidentified; emerged after benzene.  
b) Unidentified; emerged after MCH.  
c) Unidentified; emerged after toluene.  
d) Catalyst almost completely deactivated after 10 minutes.

Table LPO. DEHYDROXYMATION OF MCH OVER SXILL 45

Pressure: 1 atm Catalyst Volume: 7 ml  
Block Temperature: 642 °F Reaction Time: 30 Minutes  
Catalyst No.: 10280-45

Run Number: 11215-	93	94-1	94-2	100-1	100-2	101
LHSV	5	15	30	50	80 <sup>a)</sup>	100 <sup>a)</sup>
Catalyst Bed Profile, °F	790-88 824-22 835-33 840-38	671-67 714-07 734-48 795-92	630-22 635-28 551-44 650-73	619 610 617 633	626-30 608 608 617	635-42 603-19 608 614
Reactor Wall Temperature, °F	850	774-70	725-20	705	698	673-91
OT max	0	0	0	0	4 <sup>a)</sup>	7 <sup>a)</sup>
Product Analysis, %						
Benzene	9.9	0.9	0.1	0.0	0.0	0.1
MCH	1.1	0.5	19.3	41.2	54.6	60.3
Toluene	89.0	98.6	80.6	58.7	45.4	39.5
MCH Conversion, %	98.9	999.5	80.7	58.8	47.7	41.8

a) Back pressure was about 15 psig during this run.



Table 121. DEACTIVATION OF MCH OVER UOP-42

Pressure: 1 atm  
Block Temperature: 542°Y Reaction Time: 30 Minutes

Run Number: 11013	94-1	94-2	94
LEIV	5	15	30
Catalyst Bed Profile, °Y	770-72 799-90 815-19 831-31	709-11 626-82 716-09 739-45	752-838 540-878 671-836 682-896
Reactor Wall Temperature, °Y	815-30	775	768-858
ΔT max	4	2	168 <sup>d)</sup>
Product Analysis, %			
Benzene	3.5	0.5	0
U <sub>1</sub> <sup>a)</sup>	0.0	0.0	0.3
MCH	1.3	6.1	86.7
U <sub>2</sub> <sup>b)</sup>	0.0	0.0	0.5
Toluene	95.0	93.3	12.5
U <sub>3</sub> <sup>c)</sup>	0.2	0.0	2.0
MCH Conversion, %	98.7	95.9	15.3

- a) Unidentified; emerged after benzene.  
b) Unidentified; emerged after MCH.  
c) Unidentified; emerged after toluene.  
d) ΔT max = 162 after 17 minutes; catalyst completely deactivated at end of run.

Table 122. HYDROGENATION OF MCH OVER STANDARD CATALYST

Pressure: 1 atm Catalyst Volume: 7 ml  
 Block Temperature: 842°F Reaction Time: 30 Minutes  
 Catalyst No.: 9874-T

Run Number: 11011-	73	72-1	72-2	00
LRSP	5	15	30	50
Catalyst Bed Profile, °F	779 775 850 837-35	714-16 743 765 788-86	725-76 725-59 734-54 742-59	815-37 797-833 783-833 777-830
Reactor Wall Temperature, °F	826	792	793-97	815-34
of wall	0	2	50	52 <sup>b)</sup>
Product Analysis, %				
Benzene	1.7	0.4	0.2	0.1
MCH	1.7	19.9	62.8	93.1
Toluene	97.2	79.7	37.0	5.6
U <sub>1</sub> <sup>a)</sup>	0.0	0.0	0.0	1.2
MCH Conversion, %	98.9	80.1	37.2	6.9

a) Unidentified; emerged after toluene.

b) Catalyst almost completely deactivated at end of run.

Table 123. DEHYDROGENATION OF MCH OVER SHELL 108

Pressure: 1 atm Catalyst Volume: 7 ml  
Block Temperature: 842°F Reaction Time: 30 Minutes  
Catalyst Number: 10280-108

Run Number: 11018-	88-1	89-1	89-2	90-1	90-2	91
LESV	5	15	30	50	80	100
Catalyst Bed Profile, °F	774-72 817-13 831 837-35	653 639 752-30 775	635-39 637 657-55 687-84	642-53 630-33 637-39 658	669-75 635-39 635-37 648	687-94 644-50 639-42 646-48
Reactor Wall Temperature, °F	826-24	761	727-25	716-18	716	723-29
OF max	0	0	4	11	6	7
Product Analysis, %						
Benzene	2.3	0.2	0.0	0.0	0.0	0.0
MCH	1.4	4.5	30.4	50.8	63.3	68.3
Toluene	96.3	94.9	69.6	49.2	35.7	31.7
MCH Conversion, %	98.6	95.1	69.6	49.3	36.3	31.9

Table 124. DEHYDROGENATION OF MCH OVER Pd-150

Pressure: 1 atm Catalyst Volume: 7 ml  
 Block Temperature: 842°F Reaction Time: 30 Minutes

Run Number: 11018-	103-1	103-2	104-1	104-2	105
LHSV	5	15	30	50	80
Catalyst Bed Profile, °F	761-52 806-01 806-24 833	669-65 691-87 729-23 770-68	657-71 639-40 653-55 678-80	700-815 642-761 644-701 657-680	830-33 870-33 806-33 776-833
Reactor Wall Temperature, °F	822-20	763-61	727-29	725-88	824-35
ΔT max	0	0	14	115	-- <sup>d)</sup>
Product Analysis, %					
Benzene	1.2	0.0	0.0	0.0	0.0
U <sub>1</sub> <sup>a)</sup>	0.0	0.1	0.0	0.2	0.2
MCH	1.4	2.2	27.5	56.1	96.6
U <sub>2</sub> <sup>b)</sup>	0.0	0.0	0.0	0.0	0.6
Toluene	97.4	97.7	72.5	43.7	1.6
U <sub>3</sub> <sup>c)</sup>	0.0	0.0	0.0	0.0	1.0
MCH Conversion, %	98.6	97.8	72.5	43.5	3.4 <sup>d)</sup>

a) Unidentified; emerged after benzene.

b) Unidentified; emerged after MCH.

c) Unidentified; emerged after Toluene.

d) Catalyst about completely deactivated at end of run.

Table 125. DEHYDROGENATION OF MCH OVER UOP-R16K

Pressure: 1 atm  
 Block Temperature: 542°F  
 Catalyst Volume: 7 ml  
 Reaction Time: 30 minutes

Run No. 11325-	85-1	85-2	86-1	86-2	87-1	87-2
LHSV	5	15	30	50	80	100
Catalyst Bed Profile, °F	774-70 813-10 831-29 833	671 705-02 752-50 779-74	655-58 646-57 668-67 689-87	686-729 640-657 644-51 657-55	759-824 673-761 550-89 648-69 <sup>a)</sup>	824 784-822 705-761 678-720 <sup>a)</sup>
Reactor Wall Temp, °F	831	779-76	643-47	638-47	750-90	802-28
$\Delta T_{max}$ , °F	-4	-5	11	43	88 <sup>a)</sup>	31 <sup>a)</sup>
Product Analysis, %w						
Benzene	0.1	0.0	0.0	0.0	0.0	0.0
MCH	7.4	4.8	33.2	54.9	70.9	81.5
Toluene	92.4	95.2	66.8	45.1	29.0	18.0
Others <sup>b)</sup>	0.1	0.0	0.0	0.0	0.1	0.5
MCH Conversion, %w	92.6	95.2	66.8	45.1	29.1	18.5

a) Cold spot moved down the catalyst bed.

b) Emerged after MCH and after toluene.

Table 126. DEHYDROGENATION OF MCH OVER 1% Pt on  $\text{Al}_2\text{O}_3$ 

Pressure: 1 atm  
 Block Temperature: 542°F  
 Catalyst Volume: 7 ml  
 Reaction Time: 30 minutes  
 Catalyst No.: 10280-44

Run No. 11325-	81-1	81-2	82-1	82-2	83-1	83-2
LHSV	5	15	30	50	80	100
Catalyst Bed Profile, °F	776-72 817 833 837	686-82 732-27 765-61 795-92	669-76 689 711-09 741-39	685-743 687-707 696-707 718-22	776-815 729-90 716-66 720-54	819-24 806-24 786-819 772-812
Reactor Wall Temp, °F	831-28	781-76	750-52	747-61	770-810	817-831
$\Delta T_{\text{max}}$ , °F	-4	-5	+7	52	61 <sup>b)</sup>	40 <sup>b)</sup>
Product Analysis, %						
Benzene	2.1	0.3	0.1	0.1	0.1	0.1
MCH	2.2	12.8	44.5	63.7	82.4	92.5
Toluene	95.7	86.9	55.2	35.8	16.7	4.9
Others <sup>a)</sup>	0.0	0.0	0.2	0.4	0.8	2.5
MCH Conversion, %	97.8	87.2	55.5	36.3	17.6	7.5
Selectivity for Toluene, %	97.9	99.7	99.4	99.4	94.5	65.3

a) Emerged after MCH and after toluene.

b) Cold spot moved down the catalyst bed.

Table 127. DEHYDROGENATION OF MCH OVER 10860-114C CATALYST

Pressure: 1 atm  
Block Temperature: 842°F  
Catalyst Volume: 7 ml  
Reaction Time: 30 minutes

Run No. 11325-	74-1	74-2	75-1	75-2	76-1	76-2
LHSV	5	15	30	50	80	100
Catalyst Bed Profile, °F	759-61 815 833 837-35	640-35 689-82 750-43 792-84	619 632-30 660-57 691-89	621-26 621 635 658	635-37 619-21 626 644	644-50 621-23 626 637-35
Reactor Wall Temp, °F	831	770-66	730-28	718	711	709
$\Delta T_{max}$ , °F	+2	-8	-3	+5	+2	+6
Product Analysis, %w						
Benzene	5.4	0.5	0.1	0.0	0.0	0.0
MCH	2.2	1.7	26.8	47.2	59.6	65.0
Toluene	92.4	97.8	73.1	52.8	40.4	35.0
MCH Conversion, %w	97.8	98.3	73.2	52.8	40.4	35.0

Table 126. DEHYDROGENATION OF MCH OVER 10660-1148 CATALYST

Pressure: 1 atm  
 Block Temperature: 842°F  
 Catalyst Volume: 7 ml  
 Reaction Time: 30 minutes

Run No. 11325-	69	70-1	70-2	71	72-1	72-2
LHSV	5	15	30	50	80	100
Catalyst Bed Profile, °F	761-52 810-04 831-29 835-33	680-75 712-07 752-45 794-79	671-80 676 648-36 727-23	689-707 673-80 656-87 707	727-58 687-709 687-714 690-702	781-806 723-63 700-25 702-16
Reactor Wall Temp, °F	828-26	779-76	752	743-47	747-63	768-95
$\Delta T_{max}$ , °F	-9	-7	+9	18	41	40 <sup>a)</sup>
Product Analysis, %						
Benzene	0.8	0.2	0.1	0.1	0.1	0.1
MCH	0.8	10.8	44.3	62.2	73.3	81.3
Toluene	98.4	89.0	55.6	37.7	26.6	18.6
MCH Conversion, %	99.2	89.2	55.7	37.8	26.7	15.7

a) Cold spot moved down the catalyst bed.



Table 129. DEHYDROGENATION OF MCH OVER 10860-113A AND  
HOUDRY 200 SR CATALYSTS

Pressure: 1 atm  
Block Temperature: 847°F  
Catalyst Volume: 7  
Reaction Time: 30 minutes

Run No. 11325-	89-1	89-2	78-1	78-2	79
Catalyst	Houdry 200 SR		10860-113A		
LHSV	5	15	5	15	30
Catalyst Bed Profile, °F	758-56 801-799 826-24 833	705-833 700-831 741-830 770-828	763-58 804-797 824-12 831-28	725-68 741-50 761-58 784-779	824-37 797-833 774-826 770-817
Reactor Wall Temp, °F	826	783-838	824-21	788-97	820-837
$\Delta T_{max}$ , °F	-2	131	-7	43	b)
Product Analysis, %					
Benzene	1.4	0.2	1.7	3.2	0.1
MCH	1.7	59.3	4.8	35.0	88.3
Toluene	96.9	34.1	93.5	64.8	10.8
Others <sup>a)</sup>	0.0	6.2	0.0	0.0	0.8
MCH Conversion, %	98.3	40.6	95.2	65.0	11.7
Selectivity for Toluene, %	98.6	84.0	92.8	99.7	92.3

a) Emerged after MCH, after benzene, after toluene.

b) Catalyst bed temperature about that of reactor wall temperature.

### Description of the Pulse Reactor

The pulse reactor was a 1/4-in. OD stainless steel tube (No. 304) 9-1/8 in. long and 0.028 in. wall thickness. Swagelok Tees were fastened at each end and one arm of the Tee served as an injection port. A rubber septum (GLC type) was held in place by the fitting nut and the feed was injected through this septum from a syringe. A five inch length of the reactor tube was surrounded by a secondary furnace liner and the whole was heated by an electric furnace. The secondary liner had seven radial drilled holes for thermocouples, and the holes were located as shown in Figure 79. A schematic diagram of the pulse reactor is shown in Figure 77.

All lines were 1/4-in. OD stainless steel tubing (No. 304). About 28 in. of line just prior to the reactor was wrapped with heating tape and constituted a gas preheater. About 8 in. of the preheater section was filled with quartz chips (10-20 mesh size).

In the pulse reactor system the carrier gas was metered through a rotameter (Figure 77) and passed through the preheater section and into the reactor. The exit gas passed into a manifold and then into the GLC. The purpose of the manifold was to maintain the exit gas pressure slightly greater than the gas pressure in the GLC. This was done by adjusting the pressure control valve and the vent valve. The manifold was wrapped with heating tape and was maintained at 302° to 356°F. The injection port temperature was about 450°F. The pressure control and the vent valves were needle valves (Hoke No. 1315) and the GLC valve was a lever operated valve (Hoke No. 490).

To carry out an experiment the reactor was brought to temperature and the carrier gas flow rate, reactor pressure and manifold pressure were adjusted by means of the appropriate flow control valves. Then with inert gas flowing to the GLC a pulse was injected through the lower injection port and subsequently analyzed. This gave an analysis of the starting material. A pulse was then injected in the top injection port, passed over the catalyst and analyzed.

In this system the space velocity was obtained from the inert gas flow rate. Figure 79 shows the pulse reactor system with the secondary furnace liner in place; Figure 80 shows the GLC analysis system.

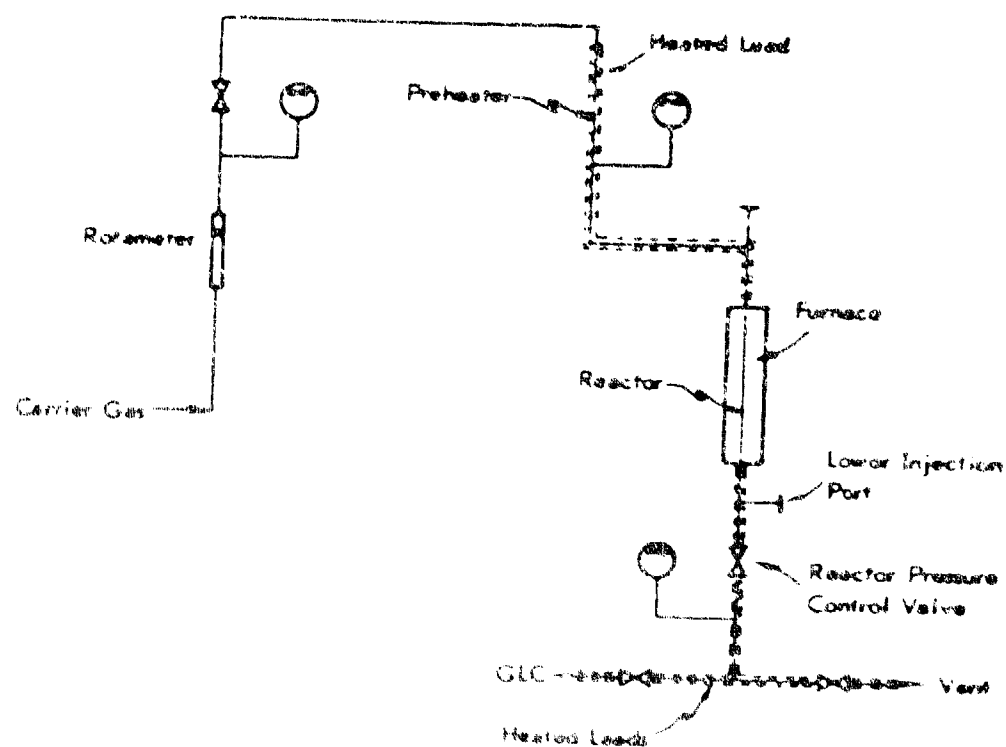


Figure 27. PULSE REACTOR SCHEMATIC

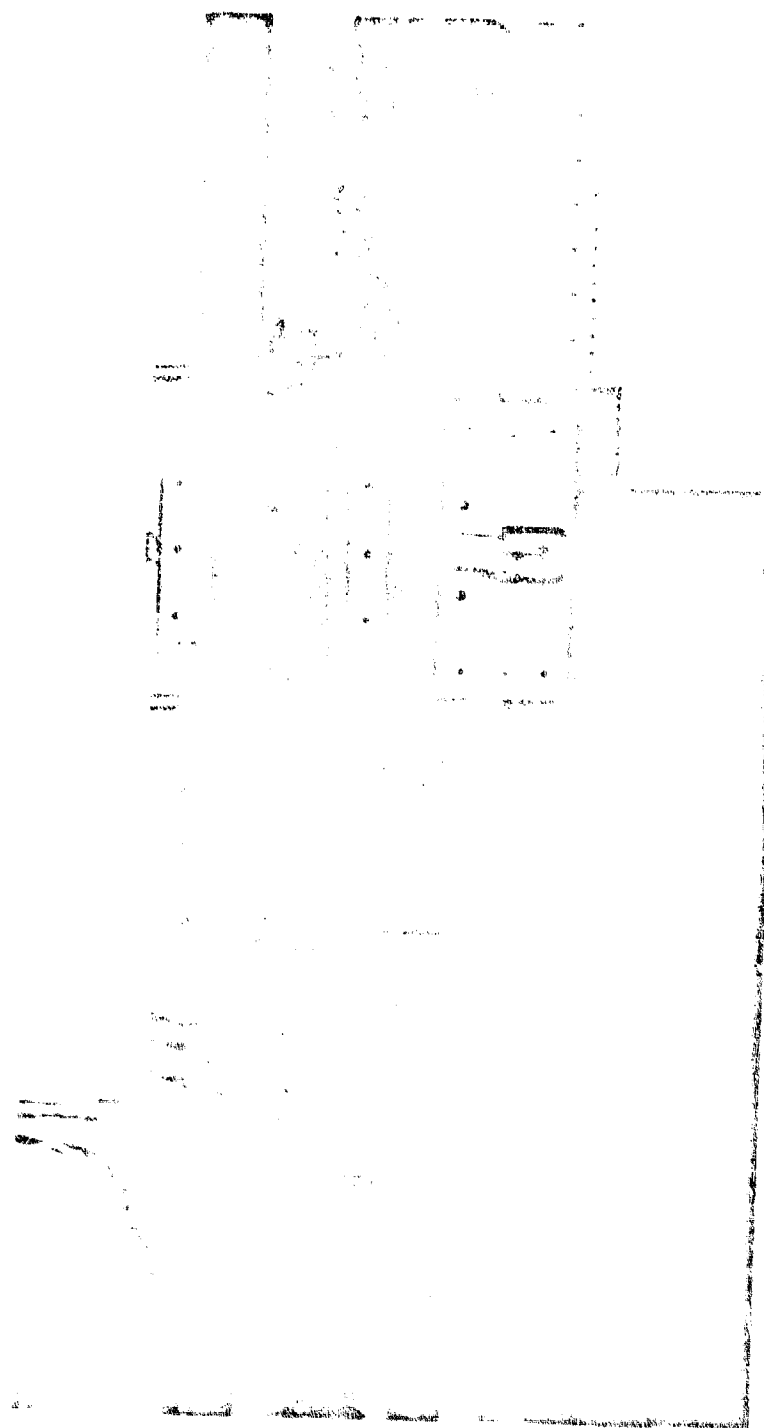


Figure 79. PULSE REACTOR SYSTEM

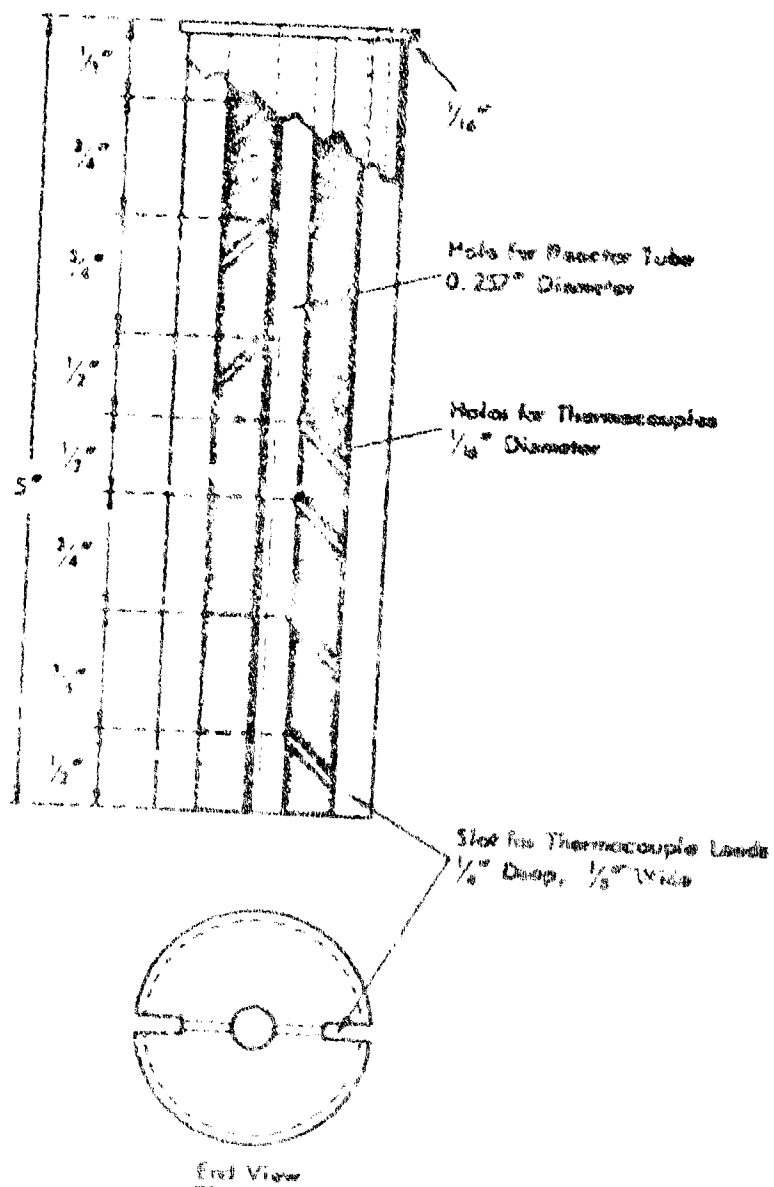


Figure 79. SECONDARY FURNACE LINER FOR PULSE REACTOR

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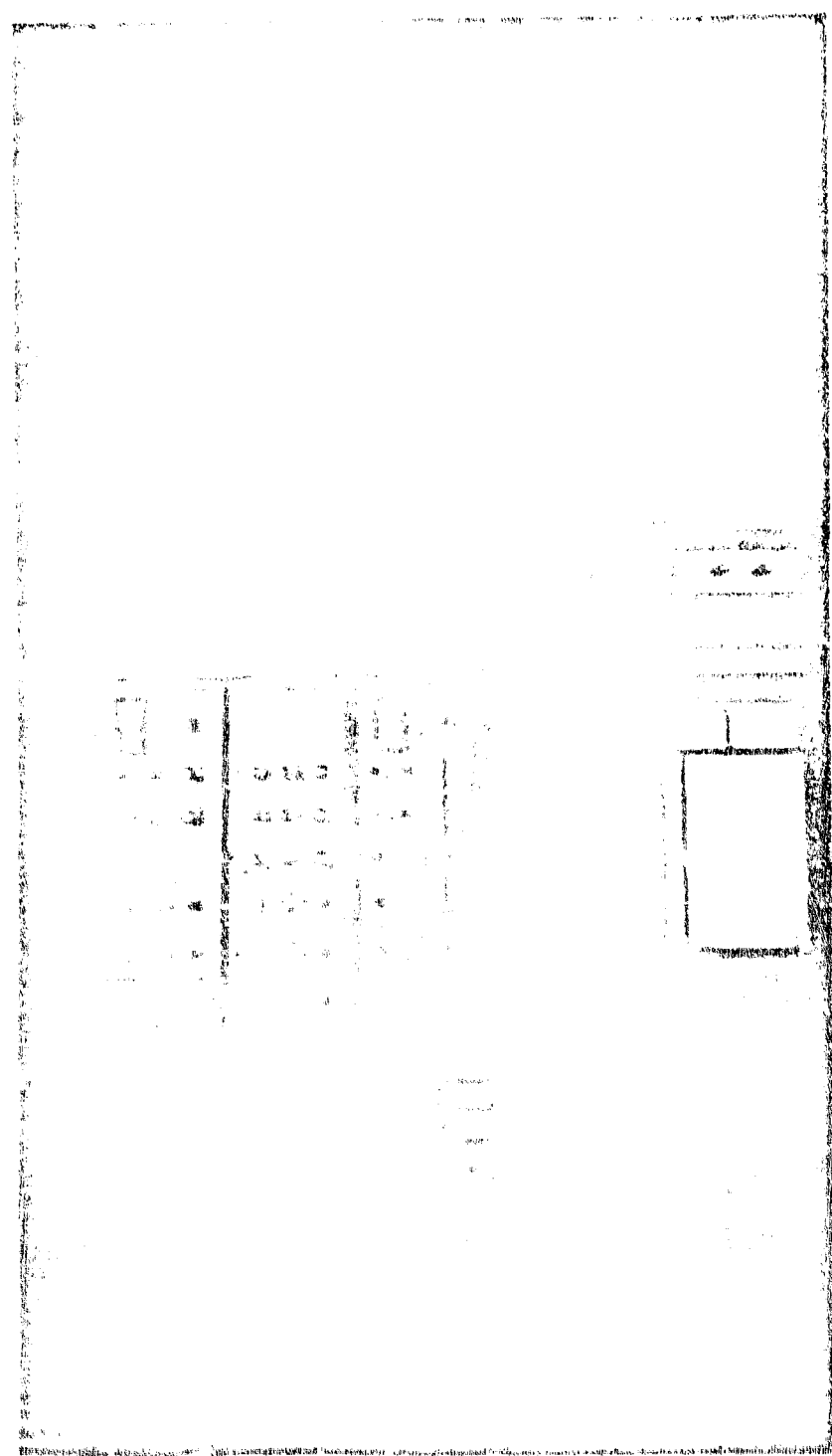


Figure 80. GIC ANALYSIS SYSTEM

### Description of the 1/4-in. OD Flow Reactor

In order to test candidate fuels that are in short supply one section of our laboratory dual reactor system was modified in the following manner, so that 1/4-in. OD reactor tubes could be used.

In our laboratory reactor system the furnace is 26 in. overall length and contains four heating elements of lengths 4", 8", 8", 4" located from top to bottom in that order. The outer shell of the furnace extends one inch beyond the top and bottom of the heating elements. The furnace consists of two hinged halves and opens lengthwise. Each half contains a heavy Meehanite liner with a groove down the center to hold the reactor tube. When closed the grooves form an opening 7/8 inch in diameter.

To modify the apparatus, a secondary furnace liner was fabricated from a 7/8-in. stainless steel rod (No. 416), 13 inches long. A 0.257-in. diameter hole was drilled down the center to accommodate a 1/4-in. OD reactor tube. Seven holes were drilled radially from the outside to the center hole in which thermocouples were cemented. The thermocouples were 1-1/2 inches apart and the top couple was 1-1/2 inches from the top of the liner. The thermocouples were situated so that they just touched the reactor wall. This secondary liner was placed in the Meehanite liners at the very bottom of the furnace and extended one inch below the bottom heating element. Figure 81 shows the construction of the secondary liner and its position in the furnace.

The reactor was a stainless steel tube (No. 304) 30 inches long, 1/4-inch OD with 0.035" wall thickness. Reaction was carried out in the lower part of the tube and the top part served as a feed preheater. The reactor was furnace-heated and a 13" long secondary furnace liner surrounded the reactor tube at the reaction zone. Figure 81 shows the secondary furnace liner and its position in the furnace.

The reactor wall temperature was measured at seven points along the tube. The points were 1-1/2 inches apart and the top point was one inch below the top of the secondary liner (Figure 81). The temperature of the reactor wall varied down the tube and Figure 82 shows the temperature variation for a furnace block temperature of 1202°F.

The maximum reaction rate will occur in the region of maximum temperature. Presumably the rate in that portion of the tube whose temperature was 18°F (10°C) or more below the maximum temperature, did not contribute appreciably to the overall rate. Thus the "effective" volume of the tube was that portion of the tube whose temperature was within 18°F of the maximum wall temperature, and whose volume was determined from a plot such as Figure 82. The "effective" reactor temperature was taken as 9°F below the maximum temperature.

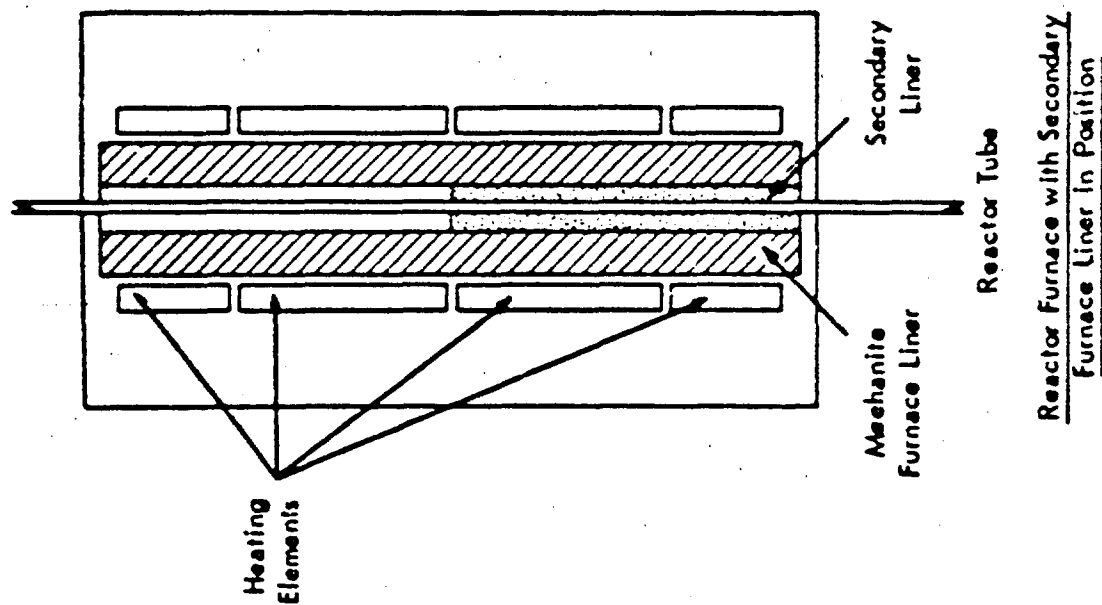
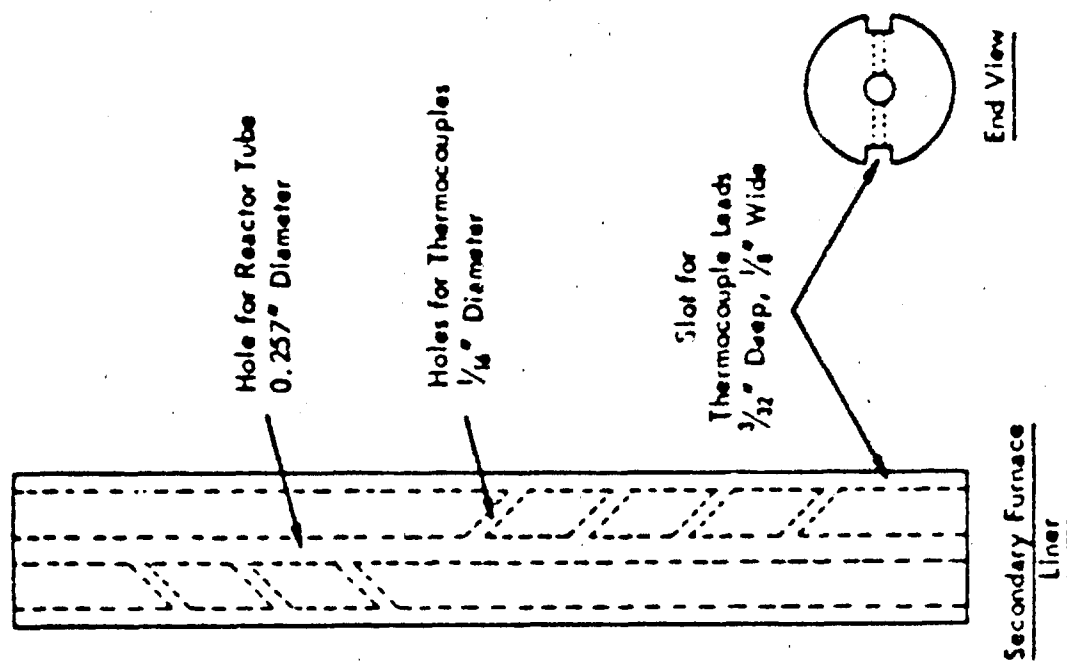


Figure 81. SECONDARY FURNACE LINER FOR 1/4" OD REACTOR TUBE



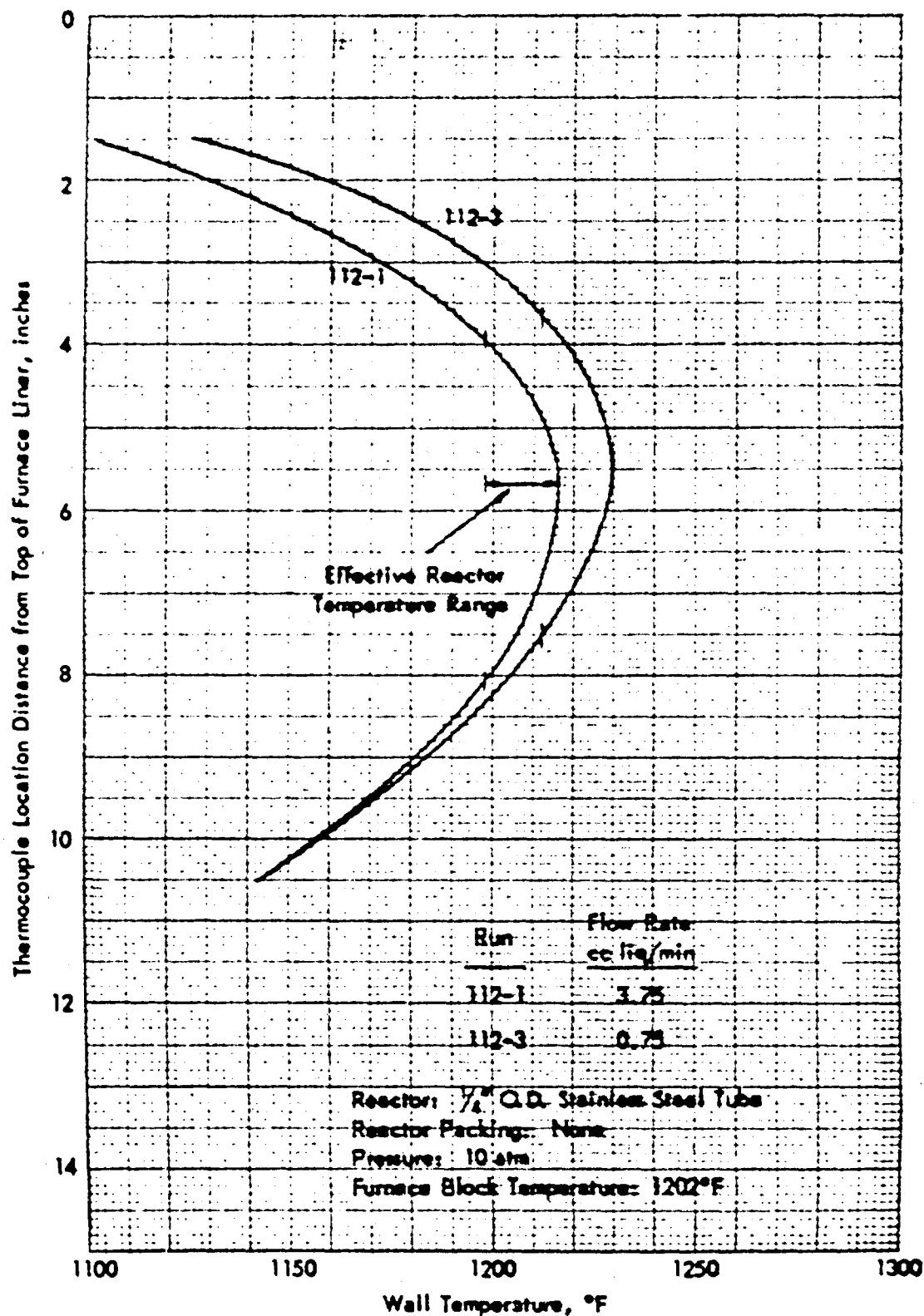


Figure 82. REACTOR TEMPERATURE PROFILE

Micro Catalyst Test Reactor Data

The micro catalyst test reactor (MCTR) and the operational techniques used for screening candidate catalysts have been described in the Appendix of the last Annual Report.<sup>16)</sup> No further changes have been made. Catalysts are tested with MCH at LHSV 100 and 662, 752 and 842°F, at 10 atm pressure without added hydrogen. Figures 87 through 89, of reference 18 show the apparatus in detail, except for the changes noted in reference 16. It has been found that more consistent results are obtained if a fresh loading of the reference catalyst 9874-139 is made each week as a base point for calibration, rather than using the same reference catalyst tube over and over again, since the activity gradually declines. Also prepared catalysts have been rescreened to 10-20 mesh to remove fines after impregnation and drying of the supports, and this gives more reproducible results.

Table 150. MEH DEHYDROGENATION WITH VARIOUS CATALYSTS IN MICR: RANS 677-813

Period: June-August 1968  
 Condition: 10 atm pressure; catalysts reduced in  $H_2$  for 20 minutes at  $T_{96}^{\circ}F$ ; GLC samples normally taken at 3-, 8-, and 13-minute operation at each block temperature.  
 Catalyst Volume: 0.9 ml catalyst diluted with 1.1 ml quartz chips; LHSV 100 (catalyst and quartz particles 10-20 mesh unless otherwise noted).

Run No.	Temperature, $^{\circ}C$	Catalyst		Gas Composition, %		
		Description	Weight	$H_2$	$CH_4$	$C_2H_6$
678	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
679	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
680	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
681	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
682	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
683	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
684	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
685	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
686	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
687	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
688	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
689	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
690	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
691	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
692	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
693	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
694	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
695	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
696	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
697	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
698	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
699	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
700	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
701	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
702	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
703	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
704	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
705	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
706	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
707	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
708	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
709	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
710	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
711	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
712	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0
713	120	15 Pt/100 20 mesh quartz (not)	0.907	100, 0.0, 0.0	0.0, 0.0, 0.0	0.0, 0.0, 0.0

(Continued)

~~TOP SECRET (S-100) FOR INFORMATION ONLY - NO DISSEMINATION~~  
~~IN DATE: 1964-10-01~~

[illegible]



Date		Description		Amount		Balance	
1917	Jan 1	Balance forward					
1917	Jan 15	By Cash	100.00		100.00		
1917	Jan 20	To Cash	50.00		150.00		
1917	Jan 25	By Cash	25.00		175.00		
1917	Jan 30	To Cash	75.00		250.00		
1917	Feb 5	By Cash	100.00		350.00		
1917	Feb 10	To Cash	150.00		500.00		
1917	Feb 15	By Cash	200.00		700.00		
1917	Feb 20	To Cash	300.00		1000.00		
1917	Feb 25	By Cash	400.00		1400.00		
1917	Feb 28	To Cash	500.00		1900.00		
1917	Mar 5	By Cash	600.00		2500.00		
1917	Mar 10	To Cash	700.00		3200.00		
1917	Mar 15	By Cash	800.00		4000.00		
1917	Mar 20	To Cash	900.00		4900.00		
1917	Mar 25	By Cash	1000.00		5900.00		
1917	Mar 30	To Cash	1100.00		7000.00		
1917	Apr 5	By Cash	1200.00		8200.00		
1917	Apr 10	To Cash	1300.00		9500.00		
1917	Apr 15	By Cash	1400.00		10900.00		
1917	Apr 20	To Cash	1500.00		12400.00		
1917	Apr 25	By Cash	1600.00		14000.00		
1917	Apr 30	To Cash	1700.00		15700.00		
1917	May 5	By Cash	1800.00		17500.00		
1917	May 10	To Cash	1900.00		19400.00		
1917	May 15	By Cash	2000.00		21400.00		
1917	May 20	To Cash	2100.00		23500.00		
1917	May 25	By Cash	2200.00		25700.00		
1917	May 30	To Cash	2300.00		28000.00		
1917	Jun 5	By Cash	2400.00		30400.00		
1917	Jun 10	To Cash	2500.00		32900.00		
1917	Jun 15	By Cash	2600.00		35500.00		
1917	Jun 20	To Cash	2700.00		38200.00		
1917	Jun 25	By Cash	2800.00		41000.00		
1917	Jun 30	To Cash	2900.00		43900.00		
1917	Jul 5	By Cash	3000.00		46900.00		
1917	Jul 10	To Cash	3100.00		50000.00		
1917	Jul 15	By Cash	3200.00		53200.00		
1917	Jul 20	To Cash	3300.00		56500.00		
1917	Jul 25	By Cash	3400.00		60000.00		
1917	Jul 30	To Cash	3500.00		63500.00		
1917	Aug 5	By Cash	3600.00		67100.00		
1917	Aug 10	To Cash	3700.00		70800.00		
1917	Aug 15	By Cash	3800.00		74600.00		
1917	Aug 20	To Cash	3900.00		78500.00		
1917	Aug 25	By Cash	4000.00		82500.00		
1917	Aug 30	To Cash	4100.00		86600.00		
1917	Sep 5	By Cash	4200.00		90800.00		
1917	Sep 10	To Cash	4300.00		95100.00		
1917	Sep 15	By Cash	4400.00		99500.00		
1917	Sep 20	To Cash	4500.00		104000.00		
1917	Sep 25	By Cash	4600.00		108600.00		
1917	Sep 30	To Cash	4700.00		113300.00		
1917	Oct 5	By Cash	4800.00		118100.00		
1917	Oct 10	To Cash	4900.00		123000.00		
1917	Oct 15	By Cash	5000.00		128000.00		
1917	Oct 20	To Cash	5100.00		133100.00		
1917	Oct 25	By Cash	5200.00		138300.00		
1917	Oct 30	To Cash	5300.00		143600.00		
1917	Nov 5	By Cash	5400.00		149000.00		
1917	Nov 10	To Cash	5500.00		154500.00		
1917	Nov 15	By Cash	5600.00		160100.00		
1917	Nov 20	To Cash	5700.00		165800.00		
1917	Nov 25	By Cash	5800.00		171600.00		
1917	Nov 30	To Cash	5900.00		177500.00		
1917	Dec 5	By Cash	6000.00		183500.00		
1917	Dec 10	To Cash	6100.00		189600.00		
1917	Dec 15	By Cash	6200.00		195800.00		
1917	Dec 20	To Cash	6300.00		202100.00		
1917	Dec 25	By Cash	6400.00		208500.00		
1917	Dec 30	To Cash	6500.00		215000.00		
1917	Dec 31	Balance forward			215000.00		

1. The first step in the process is to identify the problem or issue that needs to be addressed. This involves gathering information and understanding the context of the situation.

2. Once the problem is identified, the next step is to define the objectives and goals of the project. This helps to clarify what needs to be achieved and provides a clear direction for the team.

3. The third step is to develop a plan or strategy to address the problem. This involves breaking down the problem into smaller, manageable tasks and determining the resources needed to complete each task.

4. The fourth step is to implement the plan. This involves putting the strategy into action and monitoring progress regularly to ensure that the project is on track.

5. The final step is to evaluate the results of the project. This involves assessing the outcomes against the objectives and goals and identifying any areas for improvement or further action.

Table 111. MCM DEMONSTRATION WITH VARIOUS CATALYSTS IN MCMF

Period: September-November 1968  
 Conditions: 10 atm pressure; catalysts reduced in  $H_2$  for 20 minutes at 750°F; and samples normally taken at 5, 8, and 15 minute operation at each block temperature.  
 Catalyst Volume: 0.9 ml catalyst diluted with 1.1 ml quartz chips; 100/100 (catalyst and quartz 10-20 mesh unless otherwise noted)

Run	Catalyst	Temp (°F)	MCMF			
			5 min	8 min	15 min	100/100
1	100/100	750	0.0	0.0	0.0	0.0
2	100/100	750	0.0	0.0	0.0	0.0
3	100/100	750	0.0	0.0	0.0	0.0
4	100/100	750	0.0	0.0	0.0	0.0
5	100/100	750	0.0	0.0	0.0	0.0
6	100/100	750	0.0	0.0	0.0	0.0
7	100/100	750	0.0	0.0	0.0	0.0
8	100/100	750	0.0	0.0	0.0	0.0
9	100/100	750	0.0	0.0	0.0	0.0
10	100/100	750	0.0	0.0	0.0	0.0
11	100/100	750	0.0	0.0	0.0	0.0
12	100/100	750	0.0	0.0	0.0	0.0
13	100/100	750	0.0	0.0	0.0	0.0
14	100/100	750	0.0	0.0	0.0	0.0
15	100/100	750	0.0	0.0	0.0	0.0
16	100/100	750	0.0	0.0	0.0	0.0
17	100/100	750	0.0	0.0	0.0	0.0
18	100/100	750	0.0	0.0	0.0	0.0
19	100/100	750	0.0	0.0	0.0	0.0
20	100/100	750	0.0	0.0	0.0	0.0
21	100/100	750	0.0	0.0	0.0	0.0
22	100/100	750	0.0	0.0	0.0	0.0
23	100/100	750	0.0	0.0	0.0	0.0
24	100/100	750	0.0	0.0	0.0	0.0
25	100/100	750	0.0	0.0	0.0	0.0
26	100/100	750	0.0	0.0	0.0	0.0
27	100/100	750	0.0	0.0	0.0	0.0
28	100/100	750	0.0	0.0	0.0	0.0
29	100/100	750	0.0	0.0	0.0	0.0
30	100/100	750	0.0	0.0	0.0	0.0
31	100/100	750	0.0	0.0	0.0	0.0
32	100/100	750	0.0	0.0	0.0	0.0
33	100/100	750	0.0	0.0	0.0	0.0
34	100/100	750	0.0	0.0	0.0	0.0
35	100/100	750	0.0	0.0	0.0	0.0
36	100/100	750	0.0	0.0	0.0	0.0
37	100/100	750	0.0	0.0	0.0	0.0
38	100/100	750	0.0	0.0	0.0	0.0
39	100/100	750	0.0	0.0	0.0	0.0
40	100/100	750	0.0	0.0	0.0	0.0
41	100/100	750	0.0	0.0	0.0	0.0
42	100/100	750	0.0	0.0	0.0	0.0
43	100/100	750	0.0	0.0	0.0	0.0
44	100/100	750	0.0	0.0	0.0	0.0
45	100/100	750	0.0	0.0	0.0	0.0
46	100/100	750	0.0	0.0	0.0	0.0
47	100/100	750	0.0	0.0	0.0	0.0
48	100/100	750	0.0	0.0	0.0	0.0
49	100/100	750	0.0	0.0	0.0	0.0
50	100/100	750	0.0	0.0	0.0	0.0
51	100/100	750	0.0	0.0	0.0	0.0
52	100/100	750	0.0	0.0	0.0	0.0
53	100/100	750	0.0	0.0	0.0	0.0
54	100/100	750	0.0	0.0	0.0	0.0
55	100/100	750	0.0	0.0	0.0	0.0
56	100/100	750	0.0	0.0	0.0	0.0
57	100/100	750	0.0	0.0	0.0	0.0
58	100/100	750	0.0	0.0	0.0	0.0
59	100/100	750	0.0	0.0	0.0	0.0
60	100/100	750	0.0	0.0	0.0	0.0
61	100/100	750	0.0	0.0	0.0	0.0
62	100/100	750	0.0	0.0	0.0	0.0
63	100/100	750	0.0	0.0	0.0	0.0
64	100/100	750	0.0	0.0	0.0	0.0
65	100/100	750	0.0	0.0	0.0	0.0
66	100/100	750	0.0	0.0	0.0	0.0
67	100/100	750	0.0	0.0	0.0	0.0
68	100/100	750	0.0	0.0	0.0	0.0
69	100/100	750	0.0	0.0	0.0	0.0
70	100/100	750	0.0	0.0	0.0	0.0
71	100/100	750	0.0	0.0	0.0	0.0
72	100/100	750	0.0	0.0	0.0	0.0
73	100/100	750	0.0	0.0	0.0	0.0
74	100/100	750	0.0	0.0	0.0	0.0
75	100/100	750	0.0	0.0	0.0	0.0
76	100/100	750	0.0	0.0	0.0	0.0
77	100/100	750	0.0	0.0	0.0	0.0
78	100/100	750	0.0	0.0	0.0	0.0
79	100/100	750	0.0	0.0	0.0	0.0
80	100/100	750	0.0	0.0	0.0	0.0
81	100/100	750	0.0	0.0	0.0	0.0
82	100/100	750	0.0	0.0	0.0	0.0
83	100/100	750	0.0	0.0	0.0	0.0
84	100/100	750	0.0	0.0	0.0	0.0
85	100/100	750	0.0	0.0	0.0	0.0
86	100/100	750	0.0	0.0	0.0	0.0
87	100/100	750	0.0	0.0	0.0	0.0
88	100/100	750	0.0	0.0	0.0	0.0
89	100/100	750	0.0	0.0	0.0	0.0
90	100/100	750	0.0	0.0	0.0	0.0
91	100/100	750	0.0	0.0	0.0	0.0
92	100/100	750	0.0	0.0	0.0	0.0
93	100/100	750	0.0	0.0	0.0	0.0
94	100/100	750	0.0	0.0	0.0	0.0
95	100/100	750	0.0	0.0	0.0	0.0
96	100/100	750	0.0	0.0	0.0	0.0
97	100/100	750	0.0	0.0	0.0	0.0
98	100/100	750	0.0	0.0	0.0	0.0
99	100/100	750	0.0	0.0	0.0	0.0
100	100/100	750	0.0	0.0	0.0	0.0

1. Catalysts were reduced in  $H_2$  for 20 minutes at 750°F.  
 2. Samples were normally taken at 5, 8, and 15 minute operation at each block temperature.  
 3. Catalyst and quartz 10-20 mesh unless otherwise noted.



Table III. MCH DEMETHYLOXIDATION WITH VARIOUS CATALYSTS IN MCHTP: H<sub>2</sub>O 1:1

Period: March-August, 1969  
 Conditions: 10 atm pressure; catalysts reduced in hydrogen at 726°F, GLC samples taken normally at 3-, 8- and 13-minute operation at each temperature  
 Volume: 0.9 ml catalyst diluted with 1.1 ml quartz chips (10-20 mesh). LHSV 100 with MCH

Run No.	Catalyst Number	Description	5 MCH Conversion, %			
			3 min	8 min	13 min	18 min
105	1050-001	15 Pt/50 Rhodium type 1 support 100 type 1 support on No. 212 (1 type 212 Rhodium)	0.272	21, 20, 20	50, 48, 47	50, 48, 48
107	1050-002	15 Pt/50 Rhodium type 1 support 100 type 1 support on No. 212 (1 type 212 Rhodium)	0.272	20, 20, 20	50, 51, 49	50, 51, 51
108	1050-01	1.05 cobalt type 1 support	0.284	0, 0	0, 0	3, 3, 3
109	1050-01A	1.05 cobalt type 1 support	0.278	0, 0	0	0, 3, 3
110	1050-01B	Platonic, 2, 3.00 type 1 support	0.726	29, 28, 28	48, 48, 48	49, 49, 49
111	1050-01C	Platonic, 2, 3.00 type 1 support	0.725	28, 28, 28	51, 51, 51	50, 50, 50
112	1050-01D	Platonic, 2, 3.00 type 1 support	0.725	28, 28, 28	51, 51, 51	50, 50, 50
113	1050-01E	Platonic, 2, 3.00 type 1 support	0.726	25, 25, 25	50, 47, 46	50, 50, 50
114	1050-01F	Platonic, 2, 3.00 type 1 support	0.724	0	0, 0, 0	0, 0, 0
115	1050-01G	Platonic, 2, 3.00 type 1 support	0.726	28, 28, 28	51, 52, 52	51, 51, 51
116	1050-01H	Platonic, 2, 3.00 type 1 support	0.725	14, 17, 17	50, 50, 50	51, 51, 51
117	1050-01I	15 Pt/50 Rhodium type 1 support	0.272	20, 20, 20	47, 46, 46	50, 50, 50
118	1050-01J	Platonic, 2, 3.00 type 1 support	0.726	11, 11, 11	50, 50, 50	51, 51, 51
119	1050-01K	Platonic, 2, 3.00 type 1 support	0.726	28, 28, 28	52, 50, 48	51, 51, 51
120	1050-01L	Platonic, 2, 3.00 type 1 support	0.725	0, 0	0, 0, 0	0, 0, 0
121	1050-01M	Platonic, 2, 3.00 type 1 support	0.813	33, 31, 31	50, 50, 50	51, 51, 51
122	1050-01N	Platonic, 2, 3.00 type 1 support	0.724	28, 28, 28	50, 50, 50	51, 51, 51
123	1050-01O	Platonic, 2, 3.00 type 1 support	0.725	28, 28, 28	50, 50, 50	51, 51, 51
124	1050-01P	Platonic, 2, 3.00 type 1 support	0.724	28, 28, 28	50, 50, 50	51, 51, 51
125	1050-01Q	Platonic, 2, 3.00 type 1 support	0.724	14, 14, 14	49, 49, 49	51, 51, 51
126	1050-01R	Platonic, 2, 3.00 type 1 support	0.724	28, 28, 28	48, 47, 46	51, 51, 51
127	1050-01S	Platonic, 2, 3.00 type 1 support	0.724	0, 0, 0	0, 0, 0	0, 0, 0
128	1050-01T	15 Pt/50 Rhodium type 1 support	0.271	19, 21, 21	47, 47, 46	50, 50, 50
129	1050-01U	15 Pt/50 Rhodium type 1 support	0.272	15, 14, 14	49, 49, 49	51, 51, 51
130	1050-01V	15 Pt/50 Rhodium type 1 support	0.272	20, 20, 20	48, 47, 46	50, 50, 50
131	1050-01W	Platonic, 2, 3.00 type 1 support	0.725	0, 0, 0	0, 0, 0	0, 0, 0
132	1050-01X	Platonic, 2, 3.00 type 1 support	0.722	28, 28, 28	51, 51, 51	50, 50, 50
133	1050-01Y	15 Pt/50 Rhodium type 1 support	0.272	28, 28, 28	51, 51, 51	50, 50, 50
134	1050-01Z	15 Pt/50 Rhodium type 1 support	0.272	28, 28, 28	51, 51, 51	50, 50, 50
135	1050-02A	Platonic, 2, 3.00 type 1 support	0.729	28, 27, 26	50, 50, 50	51, 51, 51
136	1050-02B	Platonic, 2, 3.00 type 1 support	0.726	120, 11, 28	50, 52, 52	50, 50, 50
137	1050-02C	Platonic, 2, 3.00 type 1 support	0.718	35, 30, 28	50, 50, 50	50, 50, 50
138	1050-02D	Platonic, 2, 3.00 type 1 support	0.726	19, 20, 19	51, 48, 46	50, 50, 50
139	1050-02E	Platonic, 2, 3.00 type 1 support	0.648	27, 24, 24	47, 46, 46	47, 47, 47
140	1050-02F	15 Pt/50 Rhodium type 1 support 100 type 1 support on No. 212 (1 type 212 Rhodium)	0.280	20, 20, 20	50, 49, 48	50, 50, 50
141	1050-02G	15 Pt/50 Rhodium type 1 support 100 type 1 support on No. 212 (1 type 212 Rhodium)	0.280	20, 20, 20	50, 50, 49	50, 50, 50
142	1050-02H	15 Pt/50 Rhodium type 1 support 100 type 1 support on No. 212 (1 type 212 Rhodium)	0.247	20, 21, 20	50, 53, 49	51, 51, 51
143	1050-02I	15 Pt/50 Rhodium type 1 support 100 type 1 support on No. 212 (1 type 212 Rhodium)	0.246	21, 23, 23	51, 48, 47	50, 50, 50
144	1050-02J	15 Pt/50 Rhodium type 1 support	0.232	0, 0, 0	46, 45, 46	49, 50, 49

(Continued)



Table 132 (Cont'd-1). NCH DEHYDROGENATION WITH VARIOUS CATALYSTS IN MECE: RUNS 906-1000

No.	Date	Description	Amount			
			Debit	Credit	Balance	Check
1	1900-01-01	Balance				
2	1900-01-05	1000.00	1000.00			
3	1900-01-10	500.00		500.00		
4	1900-01-15	250.00	250.00			
5	1900-01-20	100.00	100.00			
6	1900-01-25	75.00		75.00		
7	1900-01-30	50.00	50.00			
8	1900-02-05	25.00	25.00			
9	1900-02-10	10.00	10.00			
10	1900-02-15	5.00	5.00			
11	1900-02-20	2.50	2.50			
12	1900-02-25	1.25	1.25			
13	1900-03-01	0.62	0.62			
14	1900-03-05	0.31	0.31			
15	1900-03-10	0.15	0.15			
16	1900-03-15	0.07	0.07			
17	1900-03-20	0.04	0.04			
18	1900-03-25	0.02	0.02			
19	1900-03-30	0.01	0.01			
20	1900-04-01	0.00	0.00			
21	1900-04-05	0.00	0.00			
22	1900-04-10	0.00	0.00			
23	1900-04-15	0.00	0.00			
24	1900-04-20	0.00	0.00			
25	1900-04-25	0.00	0.00			
26	1900-04-30	0.00	0.00			
27	1900-05-01	0.00	0.00			
28	1900-05-05	0.00	0.00			
29	1900-05-10	0.00	0.00			
30	1900-05-15	0.00	0.00			
31	1900-05-20	0.00	0.00			
32	1900-05-25	0.00	0.00			
33	1900-05-30	0.00	0.00			
34	1900-06-01	0.00	0.00			
35	1900-06-05	0.00	0.00			
36	1900-06-10	0.00	0.00			
37	1900-06-15	0.00	0.00			
38	1900-06-20	0.00	0.00			
39	1900-06-25	0.00	0.00			
40	1900-06-30	0.00	0.00			
41	1900-07-01	0.00	0.00			
42	1900-07-05	0.00	0.00			
43	1900-07-10	0.00	0.00			
44	1900-07-15	0.00	0.00			
45	1900-07-20	0.00	0.00			
46	1900-07-25	0.00	0.00			
47	1900-07-30	0.00	0.00			
48	1900-08-01	0.00	0.00			
49	1900-08-05	0.00	0.00			
50	1900-08-10	0.00	0.00			
51	1900-08-15	0.00	0.00			
52	1900-08-20	0.00	0.00			
53	1900-08-25	0.00	0.00			
54	1900-08-30	0.00	0.00			
55	1900-09-01	0.00	0.00			
56	1900-09-05	0.00	0.00			
57	1900-09-10	0.00	0.00			
58	1900-09-15	0.00	0.00			
59	1900-09-20	0.00	0.00			
60	1900-09-25	0.00	0.00			
61	1900-09-30					

(b) (5) DPP, (b) (5) ACP

Table 131 (Contd-2). HIGH DEHYDROGENATION WITH VARIOUS CATALYSTS IN MCH: H<sub>2</sub>O 100:100

No.	Date	Description	Summary			
			1950	1951	1952	1953
1000	1000-1000	1000-1000	1000	1000	1000	1000
1001	1000-1001	1000-1001	1000	1000	1000	1000
1002	1000-1002	1000-1002	1000	1000	1000	1000
1003	1000-1003	1000-1003	1000	1000	1000	1000
1004	1000-1004	1000-1004	1000	1000	1000	1000
1005	1000-1005	1000-1005	1000	1000	1000	1000
1006	1000-1006	1000-1006	1000	1000	1000	1000
1007	1000-1007	1000-1007	1000	1000	1000	1000
1008	1000-1008	1000-1008	1000	1000	1000	1000
1009	1000-1009	1000-1009	1000	1000	1000	1000
1010	1000-1010	1000-1010	1000	1000	1000	1000
1011	1000-1011	1000-1011	1000	1000	1000	1000
1012	1000-1012	1000-1012	1000	1000	1000	1000
1013	1000-1013	1000-1013	1000	1000	1000	1000
1014	1000-1014	1000-1014	1000	1000	1000	1000
1015	1000-1015	1000-1015	1000	1000	1000	1000
1016	1000-1016	1000-1016	1000	1000	1000	1000
1017	1000-1017	1000-1017	1000	1000	1000	1000
1018	1000-1018	1000-1018	1000	1000	1000	1000
1019	1000-1019	1000-1019	1000	1000	1000	1000
1020	1000-1020	1000-1020	1000	1000	1000	1000
1021	1000-1021	1000-1021	1000	1000	1000	1000
1022	1000-1022	1000-1022	1000	1000	1000	1000
1023	1000-1023	1000-1023	1000	1000	1000	1000
1024	1000-1024	1000-1024	1000	1000	1000	1000
1025	1000-1025	1000-1025	1000	1000	1000	1000
1026	1000-1026	1000-1026	1000	1000	1000	1000
1027	1000-1027	1000-1027	1000	1000	1000	1000
1028	1000-1028	1000-1028	1000	1000	1000	1000
1029	1000-1029	1000-1029	1000	1000	1000	1000
1030	1000-1030	1000-1030	1000	1000	1000	1000
1031	1000-1031	1000-1031	1000	1000	1000	1000
1032	1000-1032	1000-1032	1000	1000	1000	1000
1033	1000-1033	1000-1033	1000	1000	1000	1000
1034	1000-1034	1000-1034	1000	1000	1000	1000
1035	1000-1035	1000-1035	1000	1000	1000	1000
1036	1000-1036	1000-1036	1000	1000	1000	1000
1037	1000-1037	1000-1037	1000	1000	1000	1000
1038	1000-1038	1000-1038	1000	1000	1000	1000
1039	1000-1039	1000-1039	1000	1000	1000	1000
1040	1000-1040	1000-1040	1000	1000	1000	1000
1041	1000-1041	1000-1041	1000	1000	1000	1000
1042	1000-1042	1000-1042	1000	1000	1000	1000
1043	1000-1043	1000-1043	1000	1000	1000	1000
1044	1000-1044	1000-1044	1000	1000	1000	1000
1045	1000-1045	1000-1045	1000	1000	1000	

(Overlapped)

Table 111 (Cont'd-3). MCH DEHYDROGENATION WITH VARIOUS  
CATALYSTS IN MCH: PUMP 400-1000

Run No.	Catalyst Number	Description	5-HR Conversion, %			
			75°C	100°C	125°C	150°C
1070	1080-109	15 Pt/200 0-6 type Al <sub>2</sub> O <sub>3</sub> (ref.)	0.434	27. 24. 25	50. 46. 48	79. 77. 71
1080	1080-109	15 Pt/type 1 support	0.497	25. 24. 25	49. 47. 48	73. 73. 70
1091	1080-109	15 Pt/200 0-6 type Al <sub>2</sub> O <sub>3</sub> (ref.)	0.433	27. 25. 26	51. 47. 46	73. 71. 70
1092	1080-109	25 Pt/100 type 1 support 100 type 1 support 200 type 6 binder	0.382	24. 23. 25	38. 30. 30	62. 51. 50
1093	1080-109	25 Pt/100 type 1 support 100 type 1 support 200 type 6 binder	0.438	24. 24. 25	38. 36. 33	65. 60. 58
1094	1080-109	25 Pt/100 type 1 support 100 type 1 support 200 type 6 binder	0.480	25. 24. 27	49. 47. 46	64. 61. 58
1095	1080-109	15 Pt/100 type 1 support 100 type 1 support 200 type 6 binder	0.445	24. 24. 25	51. 48. 49	72. 70. 70
1096	1080-109	15 Pt/200 0-6 type Al <sub>2</sub> O <sub>3</sub> (ref.)	0.432	27. 26. 25	51. 48. 46	73. 73. 71
1097	1080-109	100 Pt/100 type 1 support (ref.) 100 type 1 support 200 type 6 binder Tube No. 25		25. (25). 25	79. 58. 57	79. 78. 70
1098		Same tube No. 25 filled with 200 Pt/100 100 metalized alum	0.433			
1099	1080-1113	15 Pt/100 type 1 support 100 type 1 support 200 type 6 binder	0.508	25. 25. 25	55. 51. 51	80. 81. 82
1100	1080-1113	25 Pt/100 type 1 support 100 type 1 support 200 type 6 binder	0.548	29. 27. 27	46. 46. 46	82. 83. 79
1101	1080-1112	25 Pt/100 type 1 support 100 type 1 support 200 type 6 binder	0.465	27. 25. 25	49. 47. 48	79. 76. 74
1102	1080-1112	25 Pt/100 type 1 support 100 type 1 support 200 type 6 binder	0.470	28. 27. 29	55. 55. 53	80. 81. 83
1103	1080-1112	15 Pt/100 type 1 support 100 type 1 support 200 type 6 binder	0.538	34. 30. 29	58. 55. 55	84. 81. 83
1104	1080-1110	25 Pt/100 type 1 support 100 type 1 support 200 type 6 binder	0.584	29. 26. 27	46. 45. 45	83. 83. 78
1105	1080-1110	15 Pt/100 type 1 support 100 type 1 support 200 type 6 binder	0.585	26. 26. 27	47. 46. 46	81. 82. 80
1106	1080-1111	25 Pt/100 type 1 support 100 type 1 support 200 type 6 binder	0.588	25. 24. 25	58. 50. 50	79. 76. 78
1107	1080-7	Tube No. 25 coated with: 5.00 Pt/100 type 1 support 100 type 1 support 200 5.0g		25. 25. 28	79. 58. 57	79. 78. 78
1108	1080-7	Tube No. 25 coated with: 5.00 Pt/100 type 1 support 100 type 1 support 200 5.0g		17. 26. 24	79. 59. 58	80. 81. 81
1109	1080-7	Tube No. 25 coated with: 5.00 Pt/100 type 1 support 100 type 1 support 200 5.0g After exp. 1097 Catalyst chamber filled with usual charge of reference catalyst	0.438	25. 25. 26	63. 61. 60	88. 79. 82
1110	1080-109	15 Pt/200 0-6 type Al <sub>2</sub> O <sub>3</sub> (ref.)	0.432	27. 26. 26	49. 47. 46	73. 71. 70
1111	1080-1114	15 Pt/100 type 1 support	0.751	24. 26. 28	27. 33. 33	88. 86. 86
1112	1080-1114	15 Pt/100 type 1 support	0.460	27. 27. 25	49. 46. 48	80. 83. 83

- a. Reacted numbers 5 hours; unreacted numbers 5 hours of 700 to 1000 and 1000.
- b. Heated in air at 1100°F before reduction in H<sub>2</sub>.
- c. Support with 5.00 metal dried at 700°F before impregnation with the usual metal.
- d. Support with 5.00 metal dried at 1200°F in air.
- e. 1000 H<sub>2</sub>.
- f. Metal removed from the support.
- g. Different heteropoly acid than used for comparable catalyst.
- h. Heated in air at 1200°F.
- i. Compared with No. 700.
- j. Heated in air at 1200°F.
- k. Heated in air at 1200°F.
- l. Heated with ethylene diamine.
- m. Oxidized metal, neutralized.
- n. Heated in air at 1200°F.
- o. Name of Run No. 730 (Table 1).
- p. Same as filled.
- q. Catalyst used in Run 730.
- r. 5.0g impregnation.
- s. Four separate impregnations, first three thermal decomposition at 700°F, after each impregnation.

Measurement of Deposits on Coker Tubes With Nuclear Radiation<sup>a)</sup>

Presented here is a summary to date of the results and thoughts that have gone into the application of nuclear radiation as a tool for the evaluation of coker tube deposits. Covered are the general principles, electron scattering theory, preliminary experiments, trial apparatus and results, and proposed permanent instrument design. Electron backscatter appears to be the most promising approach and is the method of primary concern in the work presented here.

Thin Film Measurement With Nuclear Radiation General Principles

Thin film measurement with nuclear radiation can be accomplished either by transmission or scatter of the radiation. The problem is to select the best type and energy of radiation and guidelines to such selection that are available.<sup>56)(57)</sup> The deposits of interest have a surface density in the neighborhood of  $10^{-5}$  g/cm<sup>2</sup> equivalent to an air path of only .01 cm. This implies an arrangement based on scattering rather than absorption and the probable need of vacuum operation. Possible types of radiation applicable in the present case are summarized in the table below.

Table 13a. METHODS OF UTILIZING NUCLEAR RADIATION

Type	Source Radiation	Detected Radiation	Operation	Remarks
1	$\alpha$	$\alpha$	alpha backscatter	Low scattering coefficient, requires very high intensity source.
2	$\alpha$	x	x-ray fluorescence in coker tube	Possible method. Efficiency very dependent on tube metal.
3	$\beta$	$\beta$	beta backscatter	Preferred method.
4	$\beta$	x	x-ray fluorescence	Similar to method 2.
5	x	x	x-ray backscatter	Low efficiency.
6	x	x	x-ray fluorescence	Low efficiency.

The conclusion from this list is that electron scattering is preferred but other factors, not listed, also lead to the same conclusion. Should an alternative method be considered for investigation the use of fluorescence from alpha bombardment is probably the most promising. A transmission type of measurement is possible if radioactivity is introduced, by plating for example, onto the coker tube. This method, suggested by H. Siegel, is

a) Acknowledgment is made to Dr. R. Curtis of the Analytical Department for this work.

preferable from the point of view of measurement to any of those listed above, but the handling of radioactive tubes is a sufficient deterrent to exclude the method.

#### Electron Backscatter Theory

An approximate description of the relative backscattered electron flux to be expected from a coating of thickness  $x$  ( $\text{g}/\text{cm}^2$ ) on a base of effectively infinite thickness is given by Tittle<sup>38</sup> as:

$$\frac{I}{I_0} = \frac{\beta_1}{\mu_1 + \lambda_1} \left[ 1 - e^{-(\mu_1 + \lambda_1)x} \right] + \frac{\beta_2}{\mu_1 + \lambda_2} e^{-(\mu_1 + \lambda_2)x} \quad (53)$$

The constants  $\mu$ ,  $\lambda$ , and  $\beta$  depend upon the materials involved and the maximum beta energy, subscripts 1 and 2 referring to the coating and base respectively, and 3 to properties of both. From relations given by Tittle equation (53) in the approximation of small  $x$  can be expressed as:

$$\begin{aligned} \frac{I}{I_0} = & \left( 1 - e^{-Z_2/40} \right) + \frac{35x}{E^{1.16}} \left[ \left( \frac{Z_1}{A_1} \right) \left( \frac{331 + Z_1}{106 + Z_1} \right) \left( 1 - e^{-Z_1/40} \right) - \right. \\ & \left. \left( \frac{Z_1}{A_1} + \frac{50Z_1 + 0.31}{106 + Z_2} \right) \left( 1 - e^{-Z_2/40} \right) \right] \end{aligned} \quad (54)$$

in which  $Z_1$  and  $A_1$  are the atomic number and atomic mass of the coating,  $Z_2$  the atomic number of the substrate, and  $E$  is the maximum beta energy in Mev. Approximating the deposit composition by  $Z_1/A_1 = 0.56$  and  $Z_1 = 5.9$  on an aluminum ( $Z_2 = 13$ ) base gives

$$\frac{I}{I_0} = 0.28 - \frac{4.4x}{E^{1.16}} \quad (55)$$

as the ratio of scattered to incident flux. The statistics of counting and the general level of instrumental variables is such that it is not practical to measure a change  $dI/I_0$  of much less than 1%. Equation (55) then predicts a maximum energy of 1.1 Kev in order to detect a thickness change  $dx$  of  $10^{-6}$  cm, assuming unit density for the deposit. Some idea of the range of thickness measurable with this energy is obtained by equating (55) to zero with the result  $x = 3.10^{-5}$   $\text{g}/\text{cm}^2$ . This prediction indicates the need of a very low energy source though perhaps not as low as 1 Kev if the expected range of thickness (up to  $10^{-6}$  cm) is to be covered. Possible sources that are available are listed in Table 135. None of these sources is as low in energy as might be desired, but the least energetic sources should provide a useable compromise since even  $^{24}\text{C}$  is capable of showing some response to the heavier deposits.

Table 135. LOW ENERGY BETA SOURCES

Isotope	Half Life, Years	E <sub>max.</sub> , Kev	Range in Air, cm
<sup>210</sup> Pb	21	17	0.4
<sup>3</sup> H	12	19	0.5
<sup>63</sup> Ni	85	67	5.5
<sup>14</sup> C	5700	155	26.0

Though not immediately apparent, equation (54) dictates that the replacement of aluminum with any metal of higher atomic number will produce an increased response to a given deposit. The method depends on the difference, primarily in atomic number, between coating and base. This difference is not large with aluminum so that a successful measurement in this case assures success with heavier metals.

#### Preliminary Experiments

Initial tests were aimed at answering three questions: whether operation without a vacuum was feasible, whether to minimize absorption a windowless flow counter was practical, and what magnitude of response would be observed in practice. To this end a small counter was constructed (courtesy A. Telfer) from one inch brass tubing with a wedge shaped end opening approximately one mm wide. As a source <sup>63</sup>Ni, having a reasonable penetration in air, was utilized in the form of the chloride adsorbed on a strip of filter paper mounted near the counter entrance. Tests were made using a 3/16" aluminum rod covered with various thicknesses of mylar film and mounted 1/2 cm from the counter.

This arrangement was unsatisfactory in several respects. One difficulty, not unexpected, was a large dependence of count rate upon counter gas flow rate. This could be controlled, but drift beyond this factor occurred that could not be accounted for. Stability was sufficient to show a count difference for one mil mylar film but was wholly inadequate for the detection of deposit films. In short, it was concluded that vacuum operation, which would require a thin window counter, was necessary and that a lower energy beta source was essential.

#### Trial Apparatus and Results

The bell jar and forepump portion of a vacuum deposition apparatus were utilized in the following measurements. Feedthroughs in the base were provided for piping the flow of counter gas, the high voltage lead to the detector, and a slide fitting to which the cocker tube could be attached. This slide fitting allowed translation and rotation of the cocker tube in front of the source and detector for scanning the deposit area.

A locally constructed thin window flow counter<sup>a)</sup> was used as detector. This window is exposed via a  $3/32 \times 5/8$ " slot cut in a one inch diameter faceplate. The source was mounted directly on this face about  $1/4$ " from the slot. The source itself was a  $1/8 \times 1/4$ " section of a neutron generator target arranged with a rather simple collimator fashioned from aluminum foil. The source-to-scatterer and scatterer-to-detector distance was 2.5 cm.

The associated electronics consisted of a Baird-Atomic Model 530 Spectrometer and Printer which provides the necessary functions of high voltage supply, pulse amplifier, discriminator, counter, and timer. Originally the high voltage was carried into the vacuum to the detector through a shielded cable, but this proved unsatisfactory. Attempts at shielding and insulating were not sufficient to eliminate corona and discharge in the vacuum with resultant spurious counting. A sufficiently hard vacuum to eliminate this problem was not practical. Instead, the high voltage lead and connection to the detector were enclosed in copper tubing and arranged to remain at atmospheric pressure.

The beta source used produces as well, some x-rays. These contribute to the scattered flux which is detected and produces a background counting rate even at atmospheric pressure. As the air pressure is reduced, a point is reached where the mean free path of the scattered electrons is sufficient for them to reach the detector and be counted. It was anticipated that a maximum count rate would be reached at some pressure and that this rate would remain constant below this pressure where essentially all electrons that could be scattered toward the detector would reach it. Instead, it was found that the count rate reached a maximum and then decreased with increasing vacuum. This maximum occurred at a pressure of approximately 25 torr while below  $1/2$  torr the count rate was independent of pressure. Apparently, as the pressure is reduced there is a maximum in count rate when the coker tube, bell jar wall, and residual atmosphere all contribute to the scattering and further evacuation diminishes the air scatter more rapidly than the increased scatter from wall and coker tube. As long as the pressure remains under  $1/2$  torr this presents no difficulty.

The coker tubes used in this study are of a miniature variety, the section of interest being  $2-1/2$ " long and  $1/8$ " diameter between end sections of  $3/16$ " diameter. The deposit generally covers only a portion of the central tube section, being lightest (visually) near one shoulder, increasing in darkness toward the center of the tube and ending fairly abruptly to leave apparently bare metal beyond this point. The scattered intensity measured along such a tube is shown in Figure 83. Each point in the figure represents a 20 second count. The scatter of these points from a smooth curve is primarily due to statistical variations in counting. The rate corresponding to an uncoated tube is about 300 c/s, while on the wider tube section beyond the shoulder it is approximately 350 c/s. While the deposit in this example is a rather heavy one by visual inspection there appears to be ample sensitivity in the backscatter response. This is particularly true considering two simple improvements that could easily be introduced. The first involves better collimation of the incident beta flux to improve resolution of the

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a) A. Telfer, "A Flow-Proportional Counter for Soft x-Rays", Emeryville Technical Progress Report 271-64.

Deposit profile, the present arrangement viewing about 1.2 cm of tube at a time. The second improvement incorporates a collimator over the detector window the purpose of which is to reduce background scatter from the bell jar. This background involves both x-ray and electron scatter and amounts to about 100 c/s in the absence of a sample tube.

Results with a group of coke tubes are given in Figure 84. Again a 20 sec count was made at each point, but with the limited number of points involved a good approximation to the profile is acquired in roughly 3 min. The two curves for each sample are scans along opposite sides of the tubes. The horizontal line to the right of each curve is an adjusted rate of 300 c/s. This adjustment was necessary because of an unexplained drift, possibly arising from the electronics, which could be corrected for each scan by returning to the starting position, but which is more difficult to correct for in going from tube to tube.

Calibration points for establishing a thickness scale were obtained using mylar film and by coating a tube with films of solution cast nitrocellulose. The mylar film (1/8 mil) is approaching infinite scattering thickness which simply means that the aluminum rod no longer contributes to the scattering. At this point the count rate has dropped from 300 to 200 c/s. Reasonably consistent results were obtained with 1, 2 and 4000A of film. Applied to the five tubes in Figure 84 this gives the following results as shown in Table 136.

$$\frac{I}{I_0} = 1 - e^{-Z_2/40} + \frac{35x}{E^{1.12}} \left[ \frac{Z_1}{A_1} \left( \frac{331 + Z_1}{106 + Z_1} \right) \left( 1 - e^{-Z_1/40} \right) - \left( \frac{Z_1}{A_1} + \frac{409 Z_1^{0.31}}{106 + Z_2} \right) \left( 1 - e^{-Z_2/40} \right) \right]$$

For Aluminum base  $Z_2 = 13$   $x = \text{film thickness, g/cm}^2$

$$\frac{I}{I_0} = 0.2775 + \frac{35x}{E^{1.12}} \left[ \frac{Z_1}{A_1} \left( \frac{331 + Z_1}{106 + Z_1} \right) \left( 1 - e^{-Z_1/40} \right) - \left( \frac{Z_1}{A_1} + 4.195 Z_1^{0.31} \right) (0.2775) \right]$$

Composition	$Z_1$	$Z_1/A_1$	$Z_1^{0.31}$	$1 - e^{-Z_1/40}$	$I/I_0$	% Change <sup>a)</sup>
CH = 7.74% H	5.613	.5383	1.707	.1309	1.914	-
CH <sub>2</sub> = 14% H	5.23	.572	1.675	.1237	1.898	-2.0
CHO. <sub>25</sub> = 23.5% O	6.18	.529	1.758	.1432	1.965	+1.5
CHS. <sub>05</sub> = 11% S	6.75	.534	1.808	.1553	2.005	+3.5
CHFe. <sub>01</sub> = 4.1% Fe	6.46	.535	1.784	.1492	1.985	+2.5
CHPb. <sub>001</sub> = 1.4% Pb	6.64	.519	1.798	.1530	1.997	+3.1

a) Scaled with CH as reference.



$n$  = st. no.

$A$  = Avg at. wt. of coating

$w$  = wt. fraction

$E$  = max energy of source, MEV

$$Z_1 = Ew_1Z_1$$

$$\frac{Z_1}{A_1} = Ew_1 \frac{Z_1}{A_1}$$

Tritium = .019 (must be a very soft source)

Table 136. RESULTS ON FIVE COVER TUBES

Fuel No.	Max Count Rate Decrease	Max Film Thickness, A°	Approx Total <sup>a)</sup> Deposit, A° x cm	Visual Ratings	
				Max	Avg
1	85 c/s	5000	6000	7.0	1.3
2	70	4500	7000	6.5	1.5
3	20	1200	1500	4.0	1.6
4	50	3000	4000	6.5	1.8
5	125	8000	6000	6.5	1.0

a) Multiplication by tube circumference (1.0 cm) will give total deposit volume.

There is little agreement with the visual ratings. The scattering results are, of course, reproducible and independent of operator judgment.

The deposit composition in calculations with equation (54) was assumed to be  $CH_2O_{.25}$ . It can approach  $CHO_{.25}$  and may contain up to 5% sulfur. Again the effects of composition changes in the deposit on the scattered intensity can be estimated at least roughly from this equation. The main effect from composition changes is in the average for  $Z$ . Starting with  $CH_2O_{.25}$  and going to  $CH_2O_{.25}S_{.05}$ , that is, adding 8% sulfur is equivalent to a 2% change in apparent thickness. The dependence on sulfur or any other heavier element is large, as expected, but film measurement within 2% that does not depend on composition is still far superior to visual estimates.

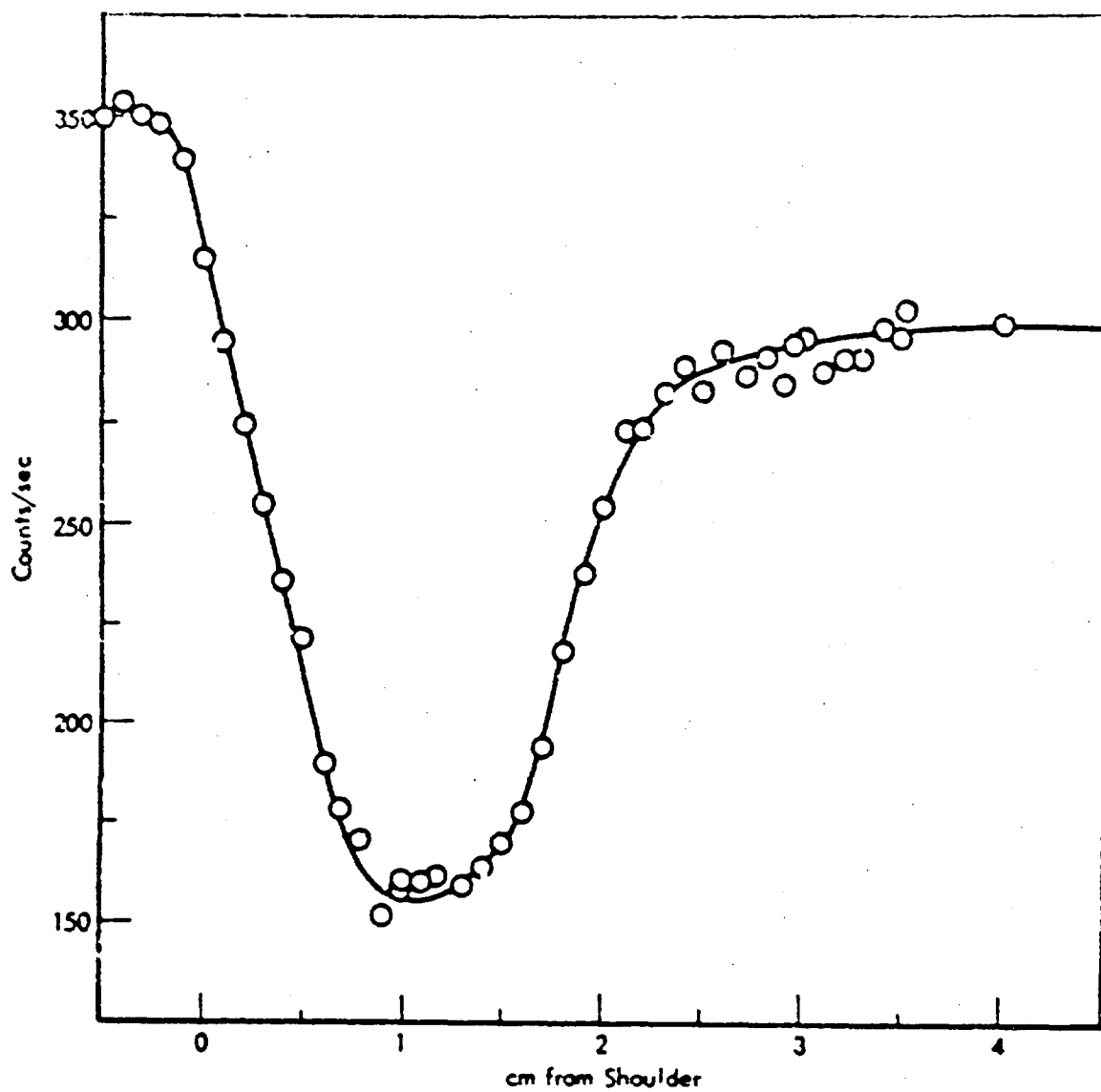


Figure 83. COKER TUBE DEPOSIT PROFILE

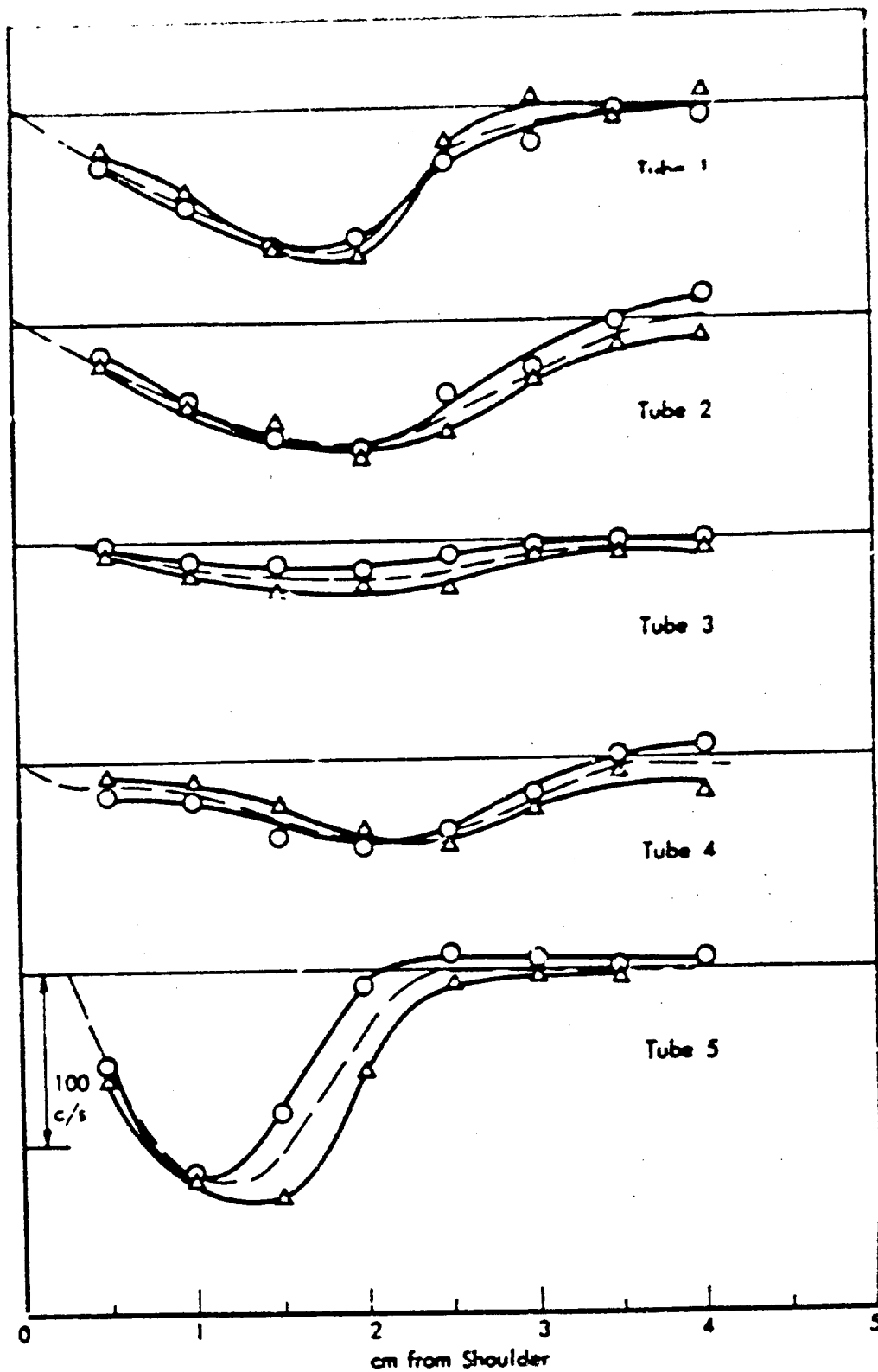


Figure 84. COMPARISON OF DEPOSIT TUBE PROFILES

### Proposed Instrument Design

The results observed above demonstrate the feasibility of electron backscatter as a means of quantitative measurement of coke tube deposits. Several conveniences can, however, be incorporated in a practical instrument. These include automatic scanning and recording of the profile and a convenient vacuum assembly.

In respect to scanning it may be convenient to integrate or average readings around the tube circumference so that a one-dimensional average along the tube length is obtained. This can be achieved in either of two ways. With a point source and detector arrangement and a ratemeter output the tube can be rotated with a period less than the time constant of the ratemeter while being translated along its length. A motor driven screw motion would achieve this. The alternative is to arrange both source and cylinder in the form of rings surrounding the rod. Actually, a triangular array would provide a sufficient approximation to a continuous ring. The main cost increase would be in respect to three detectors. The source and electronics cost increase is trivial. In this way only a linear motion need be applied to the sample tube and eccentricity is averaged out.

For the vacuum assembly it may be most convenient to adapt a commercially available apparatus involving a bell jar. But a smaller volume would allow quicker pumping down and would be more compact. This could be in the form of a tube sufficiently long for sample translation and of perhaps 3" ID. An internal motor drive would move the sample placed on a suitable carriage. Source and detector would be mounted in the mid-section wall of the tube with collimation on the source and on the detector to reduce extraneous backscatter. Electrical connections would then all be at atmospheric pressure.

TABLE 17  
LIQUID PROPERTIES OF DECALIN  
(.000 CIG. .000 TONNE)

SHELL DEPT. SHEET COMPANY

REFRACTIVE INDEX	1.462	1.462
ACENTRIC FACTOR	.299	.299
Critical Pressure	30.7 atm	21.6 MPa
Critical Temperature	413.4 C	770.2 F
Critical Compressibility Factor	.267	.267
Critical Density	.264 g/cm <sup>3</sup>	.000 g/cm <sup>3</sup>
Freezing Point	-40.8 C	-40.2 F
Normal Boiling Point	190.8 C	375.3 F
Normal Enthalpy of Vaporization	47.8 CAL/GR	127.8 BTU/LB
Heat of Combustion	10140 CAL/GR	10000 BTU/LB

TABLE 18  
LIQUID PROPERTIES OF DECALIN AT SATURATION  
(.000 CIG. .000 TONNE)

SHELL DEPT. SHEET COMPANY

TEMP. C	VAPOR PRESS. PSIA	ENTHALPY OF VAP. BTU/LB	ENTHALPY, CAL/GRAM	ENTROPY, BTU/LB-R	ENTROPY, CAL/GRAM	SPEC HEAT AT CONST P, BTU/LB-R	DENSITY, LB/CUFT	VISCOSITY, LB/FT-SEC	THERMAL COND., BTU/FT-SEC-R	PRANDTL NUMBER
-40	3.971e-04	8.903e-01	-1.227e-01	-1.000e-01	2.775e-01	9.270e-01	1.530e-01	3.740e-04	1.127e-02	1.127e-02
-30	6.167e-04	8.810e-01	-6.890e-02	-5.670e-02	3.200e-01	9.125e-01	7.571e-01	3.710e-04	1.109e-01	1.109e-01
0	1.079e-03	8.647e-01	0.000	0.000	3.490e-01	8.900e-01	4.200e-01	3.600e-04	4.302e-01	4.302e-01
10	3.400e-03	8.514e-01	7.272e-02	4.714e-02	3.707e-01	8.637e-01	2.701e-01	3.400e-04	2.803e-01	2.803e-01
20	7.751e-03	8.360e-01	1.511e-01	9.142e-02	4.270e-01	8.400e-01	1.437e-01	3.507e-04	2.803e-01	2.803e-01
30	1.500e-02	8.193e-01	2.340e-01	1.334e-01	4.340e-01	8.370e-01	1.370e-01	3.507e-04	1.401e-01	1.401e-01
40	3.120e-02	8.012e-01	3.250e-01	1.730e-01	4.600e-01	8.300e-01	1.307e-01	3.507e-04	1.307e-01	1.307e-01
100	4.375e-02	7.875e-01	4.773e-01	2.175e-01	5.215e-01	8.250e-01	7.000e-01	3.507e-04	1.107e-01	1.107e-01
120	1.200e-01	7.610e-01	5.270e-01	2.470e-01	5.310e-01	8.201e-01	6.000e-01	3.507e-04	1.107e-01	1.107e-01
140	2.400e-01	7.400e-01	6.310e-01	2.800e-01	5.400e-01	8.150e-01	5.000e-01	3.507e-04	1.107e-01	1.107e-01
160	4.500e-01	7.150e-01	7.070e-01	3.200e-01	5.500e-01	8.100e-01	4.500e-01	3.507e-04	1.107e-01	1.107e-01
180	8.000e-01	6.800e-01	8.000e-01	3.600e-01	5.600e-01	8.050e-01	4.000e-01	3.507e-04	1.107e-01	1.107e-01
200	1.300e-01	6.400e-01	9.000e-01	3.800e-01	5.600e-01	8.000e-01	3.500e-01	3.507e-04	1.107e-01	1.107e-01
220	2.100e-01	6.100e-01	1.100e-01	4.200e-01	5.700e-01	7.950e-01	3.000e-01	3.507e-04	1.107e-01	1.107e-01
240	3.200e-01	5.700e-01	1.200e-01	4.500e-01	5.700e-01	7.900e-01	2.500e-01	3.507e-04	1.107e-01	1.107e-01
260	4.600e-01	5.100e-01	1.300e-01	4.800e-01	5.700e-01	7.850e-01	2.100e-01	3.507e-04	1.107e-01	1.107e-01
280	6.500e-01	4.400e-01	1.400e-01	5.100e-01	5.700e-01	7.800e-01	1.700e-01	3.507e-04	1.107e-01	1.107e-01
300	8.900e-01	3.600e-01	1.500e-01	5.400e-01	5.700e-01	7.750e-01	1.400e-01	3.507e-04	1.107e-01	1.107e-01
320	1.12e+01	4.000e-01	1.600e-01	5.700e-01	5.700e-01	7.700e-01	1.200e-01	3.507e-04	1.107e-01	1.107e-01
340	1.420e+01	3.900e-01	1.600e-01	6.000e-01	5.700e-01	7.650e-01	1.000e-01	3.507e-04	1.107e-01	1.107e-01
360	1.700e+01	3.800e-01	1.600e-01	6.300e-01	5.700e-01	7.600e-01	8.000e-02	3.507e-04	1.107e-01	1.107e-01
380	2.100e+01	3.600e-01	1.600e-01	6.600e-01	5.700e-01	7.550e-01	7.000e-02	3.507e-04	1.107e-01	1.107e-01
400	2.300e+01	3.400e-01	1.600e-01	6.900e-01	5.700e-01	7.500e-01	6.000e-02	3.507e-04	1.107e-01	1.107e-01

TABLE 19  
LIQUID PROPERTIES OF DECALIN AT SATURATION  
(.000 CIG. .000 TONNE)

TEMP. C	VAPOR PRESS. PSIA	ENTHALPY OF VAP. BTU/LB	ENTHALPY, BTU/LB	ENTROPY, BTU/LB-R	ENTROPY, BTU/LB-R	SPEC HEAT AT CONST P, BTU/LB-R	DENSITY, LB/CUFT	VISCOSITY, LB/FT-SEC	THERMAL COND., BTU/FT-SEC-R	PRANDTL NUMBER
-40	4.700e-03	1.420e-02	-1.510e-01	-7.700e-02	2.801e-01	9.010e-01	4.400e-01	9.100e-04	1.107e-02	1.107e-02
0	1.700e-02	1.500e-02	0.000	0.000	3.270e-01	9.000e-01	1.700e-01	9.000e-04	1.107e-02	1.107e-02
10	3.500e-02	1.500e-02	1.710e-01	8.800e-02	3.610e-01	9.300e-01	8.100e-01	8.700e-04	1.107e-02	1.107e-02
100	9.871e-02	1.537e-02	9.030e-01	1.310e-01	4.200e-01	9.200e-01	4.800e-01	9.700e-04	1.107e-02	1.107e-02
120	2.717e-01	1.400e-02	9.700e-01	1.800e-01	4.400e-01	9.300e-01	2.900e-01	9.500e-04	1.107e-02	1.107e-02
140	7.300e-01	1.410e-02	8.000e-01	2.400e-01	4.900e-01	9.170e-01	2.070e-01	9.370e-04	1.107e-02	1.107e-02
160	1.070e+00	1.300e-02	1.500e-01	2.900e-01	5.300e-01	9.100e-01	1.500e-01	9.270e-04	1.107e-02	1.107e-02
180	1.600e+00	1.310e-02	1.330e-01	3.400e-01	5.600e-01	9.000e-01	1.210e-01	9.000e-04	1.107e-02	1.107e-02
200	2.200e+00	1.200e-02	1.070e-01	3.800e-01	5.700e-01	8.900e-01	9.700e-02	8.700e-04	1.107e-02	1.107e-02
220	2.810e+00	1.177e-02	1.070e-01	4.020e-01	5.700e-01	8.800e-01	8.500e-02	8.500e-04	1.107e-02	1.107e-02
240	3.400e+00	1.000e-02	2.201e-01	4.300e-01	5.700e-01	8.700e-01	7.200e-02	8.200e-04	1.107e-02	1.107e-02
260	4.000e+00	1.010e-02	2.602e-01	4.500e-01	5.700e-01	8.600e-01	6.300e-02	8.100e-04	1.107e-02	1.107e-02
280	4.600e+00	9.717e-03	2.907e-01	4.700e-01	5.700e-01	8.500e-01	5.300e-02	8.000e-04	1.107e-02	1.107e-02
300	5.300e+00	8.100e-03	3.200e-01	4.900e-01	5.700e-01	8.400e-01	4.500e-02	7.900e-04	1.107e-02	1.107e-02
320	6.100e+00	6.900e-03	3.510e-01	5.100e-01	5.700e-01	8.300e-01	3.800e-02	7.800e-04	1.107e-02	1.107e-02
340	7.000e+00	6.000e-03	3.710e-01	5.300e-01	5.700e-01	8.200e-01	3.200e-02	7.700e-04	1.107e-02	1.107e-02
360	8.000e+00	5.400e-03	4.120e-01	5.500e-01	5.700e-01	8.100e-01	2.800e-02	7.600e-04	1.107e-02	1.107e-02
380	9.100e+00	4.800e-03	4.400e-01	5.700e-01	5.700e-01	8.000e-01	2.400e-02	7.500e-04	1.107e-02	1.107e-02
400	1.000e+01	4.400e-03	4.600e-01	5.900e-01	5.700e-01	7.900e-01	2.100e-02	7.400e-04	1.107e-02	1.107e-02















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Year	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017	2018	2019	2020	2021	2022	2023	2024	2025	2026	2027	2028	2029	2030	2031	2032	2033	2034	2035	2036	2037	2038	2039	2040	2041	2042	2043	2044	2045	2046	2047	2048	2049	2050	2051	2052	2053	2054	2055	2056	2057	2058	2059	2060	2061	2062	2063	2064	2065	2066	2067	2068	2069	2070	2071	2072	2073	2074	2075	2076	2077	2078	2079	2080	2081	2082	2083	2084	2085	2086	2087	2088	2089	2090	2091	2092	2093	2094	2095	2096	2097	2098	2099	2100
1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017	2018	2019	2020	2021	2022	2023	2024	2025	2026	2027	2028	2029	2030	2031	2032	2033	2034	2035	2036	2037	2038	2039	2040	2041	2042	2043	2044	2045	2046	2047	2048	2049	2050	2051	2052	2053	2054	2055	2056	2057	2058	2059	2060	2061	2062	2063	2064	2065	2066	2067	2068	2069	2070	2071	2072	2073	2074	2075	2076	2077	2078	2079	2080	2081	2082	2083	2084	2085	2086	2087	2088	2089	2090	2091	2092	2093	2094	2095	2096	2097	2098	2099	2100	



Sl. No.	Name of the Candidate	Roll No.	Grade	Subject	Score	Remarks
1	ABHIJITH K	101	10	Maths	85	
2	ADARSH K	102	10	Maths	78	
3	ADITHYAN K	103	10	Maths	92	
4	ADITHYAN K	104	10	Maths	88	
5	ADITHYAN K	105	10	Maths	75	
6	ADITHYAN K	106	10	Maths	82	
7	ADITHYAN K	107	10	Maths	79	
8	ADITHYAN K	108	10	Maths	86	
9	ADITHYAN K	109	10	Maths	81	
10	ADITHYAN K	110	10	Maths	84	
11	ADITHYAN K	111	10	Maths	87	
12	ADITHYAN K	112	10	Maths	83	
13	ADITHYAN K	113	10	Maths	80	
14	ADITHYAN K	114	10	Maths	85	
15	ADITHYAN K	115	10	Maths	82	
16	ADITHYAN K	116	10	Maths	86	
17	ADITHYAN K	117	10	Maths	81	
18	ADITHYAN K	118	10	Maths	84	
19	ADITHYAN K	119	10	Maths	87	
20	ADITHYAN K	120	10	Maths	83	
21	ADITHYAN K	121	10	Maths	80	
22	ADITHYAN K	122	10	Maths	85	
23	ADITHYAN K	123	10	Maths	82	
24	ADITHYAN K	124	10	Maths	86	
25	ADITHYAN K	125	10	Maths	81	
26	ADITHYAN K	126	10	Maths	84	
27	ADITHYAN K	127	10	Maths	87	
28	ADITHYAN K	128	10	Maths	83	
29	ADITHYAN K	129	10	Maths	80	
30	ADITHYAN K	130	10	Maths	85	
31	ADITHYAN K	131	10	Maths	82	
32	ADITHYAN K	132	10	Maths	86	
33	ADITHYAN K	133	10	Maths	81	
34	ADITHYAN K	134	10	Maths	84	
35	ADITHYAN K	135	10	Maths	87	
36	ADITHYAN K	136	10	Maths	83	
37	ADITHYAN K	137	10	Maths	80	
38	ADITHYAN K	138	10	Maths	85	
39	ADITHYAN K	139	10	Maths	82	
40	ADITHYAN K	140	10	Maths	86	
41	ADITHYAN K	141	10	Maths	81	
42	ADITHYAN K	142	10	Maths	84	
43	ADITHYAN K	143	10	Maths	87	
44	ADITHYAN K	144	10	Maths	83	
45	ADITHYAN K	145	10	Maths	80	
46	ADITHYAN K	146	10	Maths	85	
47	ADITHYAN K	147	10	Maths	82	
48	ADITHYAN K	148	10	Maths	86	
49	ADITHYAN K	149	10	Maths	81	
50	ADITHYAN K	150	10	Maths	84	
51	ADITHYAN K	151	10	Maths	87	
52	ADITHYAN K	152	10	Maths	83	
53	ADITHYAN K	153	10	Maths	80	
54	ADITHYAN K	154	10	Maths	85	
55	ADITHYAN K	155	10	Maths	82	
56	ADITHYAN K	156	10	Maths	86	
57	ADITHYAN K	157	10	Maths	81	
58	ADITHYAN K	158	10	Maths	84	
59	ADITHYAN K	159	10	Maths	87	
60	ADITHYAN K	160	10	Maths	83	
61	ADITHYAN K	161	10	Maths	80	
62	ADITHYAN K	162	10	Maths	85	
63	ADITHYAN K	163	10	Maths	82	

[illegible]



DATE OF RECEIPT: 10-18-67 BY: J. H. C. NO. 10-18-67

DATE: 24-03-2024

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1992年10月10日 星期三

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10/17/75 10:45 AM (10/17/75) 10:45 AM, 10/17/75

Year	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017	2018	2019	2020	2021	2022	2023	2024	2025	2026	2027	2028	2029	2030	2031	2032	2033	2034	2035	2036	2037	2038	2039	2040	2041	2042	2043	2044	2045	2046	2047	2048	2049	2050	2051	2052	2053	2054	2055	2056	2057	2058	2059	2060	2061	2062	2063	2064	2065	2066	2067	2068	2069	2070	2071	2072	2073	2074	2075	2076	2077	2078	2079	2080	2081	2082	2083	2084	2085	2086	2087	2088	2089	2090	2091	2092	2093	2094	2095	2096	2097	2098	2099	2100	2101	2102	2103	2104	2105	2106	2107	2108	2109	2110	2111	2112	2113	2114	2115	2116	2117	2118	2119	2120	2121	2122	2123	2124	2125	2126	2127	2128	2129	2130	2131	2132	2133	2134	2135	2136	2137	2138	2139	2140	2141	2142	2143	2144	2145	2146	2147	2148	2149	2150	2151	2152	2153	2154	2155	2156	2157	2158	2159	2160	2161	2162	2163	2164	2165	2166	2167	2168	2169	2170	2171	2172	2173	2174	2175	2176	2177	2178	2179	2180	2181	2182	2183	2184	2185	2186	2187	2188	2189	2190	2191	2192	2193	2194	2195	2196	2197	2198	2199	2200	2201	2202	2203	2204	2205	2206	2207	2208	2209	2210	2211	2212	2213	2214	2215	2216	2217	2218	2219	2220	2221	2222	2223	2224	2225	2226	2227	2228	2229	2230	2231	2232	2233	2234	2235	2236	2237	2238	2239	2240	2241	2242	2243	2244	2245	2246	2247	2248	2249	2250	2251	2252	2253	2254	2255	2256	2257	2258	2259	2260	2261	2262	2263	2264	2265	2266	2267	2268	2269	2270	2271	2272	2273	2274	2275	2276	2277	2278	2279	2280	2281	2282	2283	2284	2285	2286	2287	2288	2289	2290	2291	2292	2293	2294	2295	2296	2297	2298	2299	2300	2301	2302	2303	2304	2305	2306	2307	2308	2309	2310	2311	2312	2313	2314	2315	2316	2317	2318	2319	2320	2321	2322	2323	2324	2325	2326	2327	2328	2329	2330	2331	2332	2333	2334	2335	2336	2337	2338	2339	2340	2341	2342	2343	2344	2345	2346	2347	2348	2349	2350	2351	2352	2353	2354	2355	2356	2357	2358	2359	2360	2361	2362	2363	2364	2365	2366	2367	2368	2369	2370	2371	2372	2373	2374	2375	2376	2377	2378	2379	2380	2381	2382	2383	2384	2385	2386	2387	2388	2389	2390	2391	2392	2393	2394	2395	2396	2397	2398	2399	2400	2401	2402	2403	2404	2405	2406	2407	2408	2409	2410	2411	2412	2413	2414	2415	2416	2417	2418	2419	2420	2421	2422	2423	2424	2425	2426	2427	2428	2429	2430	2431	2432	2433	2434	2435	2436	2437	2438	2439	2440	2441	2442
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[illegible]

6-8 700000 700 50 900000 1 . 000 010 . 000 000

TABLE 1. SUMMARY OF THE DATA FOR THE 1970-1971 FLOODING OF THE MISSISSIPPI RIVER											
STATION		DATE		FLOODING		WATER LEVEL		WIND VELOCITY		WIND DIRECTION	
NO.	NAME	MO.	DAY	START	END	FEET	FEET	MPH	MPH	DEG	DEG
1	STATION 1	1	1	12:00	12:00	10.0	10.0	10	10	0	0
2	STATION 2	1	2	12:00	12:00	10.0	10.0	10	10	0	0
3	STATION 3	1	3	12:00	12:00	10.0	10.0	10	10	0	0
4	STATION 4	1	4	12:00	12:00	10.0	10.0	10	10	0	0
5	STATION 5	1	5	12:00	12:00	10.0	10.0	10	10	0	0
6	STATION 6	1	6	12:00	12:00	10.0	10.0	10	10	0	0
7	STATION 7	1	7	12:00	12:00	10.0	10.0	10	10	0	0
8	STATION 8	1	8	12:00	12:00	10.0	10.0	10	10	0	0
9	STATION 9	1	9	12:00	12:00	10.0	10.0	10	10	0	0
10	STATION 10	1	10	12:00	12:00	10.0	10.0	10	10	0	0
11	STATION 11	1	11	12:00	12:00	10.0	10.0	10	10	0	0
12	STATION 12	1	12	12:00	12:00	10.0	10.0	10	10	0	0
13	STATION 13	1	13	12:00	12:00	10.0	10.0	10	10	0	0
14	STATION 14	1	14	12:00	12:00	10.0	10.0	10	10	0	0
15	STATION 15	1	15	12:00	12:00	10.0	10.0	10	10	0	0
16	STATION 16	1	16	12:00	12:00	10.0	10.0	10	10	0	0
17	STATION 17	1	17	12:00	12:00	10.0	10.0	10	10	0	0
18	STATION 18	1	18	12:00	12:00	10.0	10.0	10	10	0	0
19	STATION 19	1	19	12:00	12:00	10.0	10.0	10	10	0	0
20	STATION 20	1	20	12:00	12:00	10.0	10.0	10	10	0	0
21	STATION 21	1	21	12:00	12:00	10.0	10.0	10	10	0	0
22	STATION 22	1	22	12:00	12:00	10.0	10.0	10	10	0	0
23	STATION 23	1	23	12:00	12:00	10.0	10.0	10	10	0	0
24	STATION 24	1	24	12:00	12:00	10.0	10.0	10	10	0	0
25	STATION 25	1	25	12:00	12:00	10.0	10.0	10	10	0	0
26	STATION 26	1	26	12:00	12:00	10.0	10.0	10	10	0	0
27	STATION 27	1	27	12:00	12:00	10.0	10.0	10	10	0	0
28	STATION 28	1	28	12:00	12:00	10.0	10.0	10	10	0	0
29	STATION 29	1	29	12:00	12:00	10.0	10.0	10	10	0	0
30	STATION 30	1	30	12:00	12:00	10.0	10.0	10	10	0	0
31	STATION 31	1	31	12:00	12:00	10.0	10.0	10	10	0	0
32	STATION 32	1	32	12:00	12:00	10.0	10.0	10	10	0	0
33	STATION 33	1	33	12:00	12:00	10.0	10.0	10	10	0	0
34	STATION 34	1	34	12:00	12:00	10.0	10.0	10	10	0	0
35	STATION 35	1	35	12:00	12:00	10.0	10.0	10	10	0	0

[illegible]

TABLE 1. SUMMARY OF DATA FOR THE 1970-1971 SEASON									
STATION	DATE	TIME	WIND DIRECTION	WIND SPEED (KNOTS)	WAVE PERIOD (SECONDS)	WAVE HEIGHT (FEET)	WAVE LENGTH (FEET)	WAVE ENERGY (KCAL/M <sup>2</sup> )	WAVE POWER (KCAL/M <sup>2</sup> )
1	1970-12-15	11:00	110	10	10	10	10	10	10
2	1970-12-15	11:00	110	10	10	10	10	10	10
3	1970-12-15	11:00	110	10	10	10	10	10	10
4	1970-12-15	11:00	110	10	10	10	10	10	10
5	1970-12-15	11:00	110	10	10	10	10	10	10
6	1970-12-15	11:00	110	10	10	10	10	10	10
7	1970-12-15	11:00	110	10	10	10	10	10	10
8	1970-12-15	11:00	110	10	10	10	10	10	10
9	1970-12-15	11:00	110	10	10	10	10	10	10
10	1970-12-15	11:00	110	10	10	10	10	10	10
11	1970-12-15	11:00	110	10	10	10	10	10	10
12	1970-12-15	11:00	110	10	10	10	10	10	10
13	1970-12-15	11:00	110	10	10	10	10	10	10
14	1970-12-15	11:00	110	10	10	10	10	10	10
15	1970-12-15	11:00	110	10	10	10	10	10	10
16	1970-12-15	11:00	110	10	10	10	10	10	10
17	1970-12-15	11:00	110	10	10	10	10	10	10
18	1970-12-15	11:00	110	10	10	10	10	10	10
19	1970-12-15	11:00	110	10	10	10	10	10	10
20	1970-12-15	11:00	110	10	10	10	10	10	10
21	1970-12-15	11:00	110	10	10	10	10	10	10
22	1970-12-15	11:00	110	10	10	10	10	10	10
23	1970-12-15	11:00	110	10	10	10	10	10	10
24	1970-12-15	11:00	110	10	10	10	10	10	10
25	1970-12-15	11:00	110	10	10	10	10	10	10
26	1970-12-15	11:00	110	10	10	10	10	10	10
27	1970-12-15	11:00	110	10	10	10	10	10	10
28	1970-12-15	11:00	110	10	10	10	10	10	10
29	1970-12-15	11:00	110	10	10	10	10	10	10
30	1970-12-15	11:00	110	10	10	10	10	10	10
31	1970-12-15	11:00	110	10	10	10	10	10	10
32	1970-12-15	11:00	110	10	10	10	10	10	10
33	1970-12-15	11:00	110	10	10	10	10	10	10
34	1970-12-15	11:00	110	10	10	10	10	10	10
35	1970-12-15	11:00	110	10	10	10	10	10	10
36	1970-12-15	11:00	110	10	10	10	10	10	10
37	1970-12-15	11:00	110	10	10	10	10	10	10
38	1970-12-15	11:00	110	10	10	10	10	10	10
39	1970-12-15	11:00	110	10	10	10	10	10	10
40	1970-12-15	11:00	110	10	10	10	10	10	10





150

1. The first group of people who are interested in the results of the study are the researchers themselves. They want to know if the study was successful in achieving its goals and if the data collected is reliable and valid. They also want to know if the study has contributed to the field of research and if it has any practical implications.

[illegible]

~~SECRET~~

[illegible]







[illegible][illegible][illegible]

100

[illegible][illegible][illegible][illegible]

[illegible][illegible]





1998

[illegible][illegible]







1. The first group of people who are interested in the study of the history of the United States are the people who are interested in the history of the United States.

**SECRET**

1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100
1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86														

6-00000000000000000000000000000000

**Abstract**

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[illegible]

















EXHIBIT (3-10)

Qualitative Analytical and Physical Property Data

Following are typical analytical data and calculated physical property information based on a variety of methods. Since only a limited number of samples of CHILLDYNE-H have been prepared, the consistency of the data from batch to batch has not been established. Also, pending larger scale preparations, we have not independently calculated the effect of temperature and pressure, outside the normal range, on various liquid and gaseous properties of interest. Instead we have applied corrections to the values calculated for CHILLDYNE itself, as given in the citation above, as deemed advisable.

The magnitudes of the correction factors involved are generally not large and we believe the data can be used without serious error. Improved information will be supplied as soon as it is available.

Table 1.1. Summary of Data

Physical and Chemical Properties of the Sample

	Typical Value	Test Method
Gravity, deg API	1.01	ASTM D-155
Specific Gravity	1.01	
Refractive Index, $n_D^{20}$	1.44	ASTM
Color	Water white	
Distillation temperature, °F		ASTM D-156
Initial boiling point	107	
1% Evaporated	115	
2% Evaporated	119	
5% Evaporated	122	
10% Evaporated	125	
End Point	127	
Loss, %	1.1	
Residue, %	0.9	
Sulfur, %	0.0001	ASTM D-129-47
Mercaptan sulfur, %	<0.001	ASTM D-129-47
Existent O <sub>2</sub> , mg per 100 ml	1	ASTM D-129-47
Freezing point, °F	-40(-40)	ASTM D-156-57
Boiling point, °F	127	ASTM D-156-57
Net heat of combustion		ASTM D-156-57
Btu/lb	17,400	
Btu/gal	101,300	
Aromatic content, %	<0.1	ASTM D-129-47
Cladin content, %	1	D-129-47
Copper strip corrosion	1A	ASTM D-156-57
Viscosity, cs, at 100°F	13	ASTM D-156-57
0°F	215	
-30°F	1250	
Flash point, °F	120	ASTM D-156-57
Vapor pressure, psia at 300°F	0.5	
psia at 500°F	12	
Thermal stability, 5 hr	typical	ERC Modified
Pressure change, in. Hg	1.5	Standard Gaer
Preheater deposit rating	2-1/2	
Smoke point	0.5	D-156-57
Carbon/hydrogen ratio	9.8	Calculated
Critical temperature, °F	400	API 441.2
Critical pressure, psia	110	API 441.2
Heat of formation, Btu/lb	100	
Molecular weight	150.7	

- a) Generally fluid because of low viscosity; some samples have formed crystals at temperatures of about -30°F and below.
- b) D-156-57 results only.
- c) All 50% point corrected as per API Figure M1.1.

Table 116. 114 IS PROPERTIES OF "E" ELEMENTS AT SATURATED PRESSURE

Temp. °F	Density, lb/ft <sup>3</sup>	Viscosity, a) lb/ft-hr	Thermal conductivity, Btu/ft-hr-°F	Heat capacity, Btu/lb-°F	Enthalpy, Btu/lb	Heat of vaporization, Btu/lb	Vapor pressure, psia
-60	70.8	22000.	0.048	0.217	-14.5	151.9	0.0220
0	69.5	532.	0.036	0.257	0.0	145.5	0.0000
100	66.6	35.7	0.093	0.342	29.0	140.3	0.3011
200	63.8	8.78	0.090	0.307	64.5	135.7	0.0000
300	60.9	3.12	0.065	0.452	106.4	128.4	0.0000
400	57.8	1.41	0.079	0.507	154.4	120.4	2.97
500	54.5	0.752	0.072	0.559	207.7	111.5	11.30
600	50.9	0.432	0.064	0.606	266.0	101.3	32.9
700	46.7	0.341	0.057	0.651	328.8	88.9	76.2
800	41.7	0.238	0.048	0.702	396.3	72.9	153.0
900	34.2	0.155	0.040	0.818	470.9	46.8	278.0

a) To convert to cgs multiply by  $\frac{25.8}{\text{density}}$ .



Table 1. (Continued) Performance of the AFAPL-TR-07-114

Performance of the AFAPL-TR-07-114										
Year	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999
Performance of the AFAPL-TR-07-114										
1990	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1991	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1992	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1993	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1994	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1995	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1996	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1997	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1998	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1999	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
Performance of the AFAPL-TR-07-114										
1990	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1991	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1992	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1993	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1994	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1995	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1996	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1997	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1998	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1999	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
Performance of the AFAPL-TR-07-114										
1990	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1991	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1992	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1993	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1994	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1995	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1996	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1997	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1998	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0
1999	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0

a) The performance index is the ratio of the observed performance to the target performance.









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Part III

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<b>VARIATION AND ENDOTHERMIC FUELS FOR ADVANCED ENGINE APPLICATION.</b> <b>Part III. Studies of Thermal and Catalytic Reactions, Thermal Stabilities,</b> <b>and Combustion Properties of Hydrocarbon Fuels</b>		
Final Report - June 1-30-September 1969		
A. C. Nixon, G. H. Ackerman, L. E. Faith, H. T. Henderson, A. W. Ritchie, L. B. Ryland and T. M. Chryne		
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Air Force Aero Propulsion Laboratory Wright Patterson Air Force Base, Ohio		
<p>The feasibility of utilizing hydrocarbon fuels for high speed air craft depends upon the endothermic and enthalpic capacity of the hydrocarbons, and the combustion properties of the products. At Mach 5 a supersonic combustion ram jet engine would require about 1900 Btu of cooling per pound of fuel. The enthalpic capacity of hydrocarbons can be augmented by thermal and catalytic reactions. These have been studied. The rate of thermal cracking can be accelerated by means of additives. The rate of dehydrogenation of naphthenes, a strongly endothermic reaction, can be increased by the use of improved catalysts based on platinum and similar. The stabilities of such catalysts are inversely proportional to the pore size of the support and are affected by composition. Dispersed catalysts have some advantages over bed type catalysts and some indications of possible success have been observed. About 1/3 of the 850 catalysts developed for naphthene dehydrogenation has been more active than the standard laboratory catalyst, although none is orders of magnitude better. Wall catalysts have the advantage of low pressure drop and are showing efficiency benefits also. Calculations show that diffusion limitations can be avoided if the coating thickness is no more than about 3 mils thick. Satisfactory operation with both improved bed catalysts and wall catalysts has been demonstrated in the fuel system simulator with both methylcyclohexane and Decalin. Heat transfer studies have been carried out with MCP, Decalin, HYDROLYNE-S and JP-7 fuel in small diameter test sections under heat fluxes up to <math>8 \times 10^5</math> Btu per hour per square foot. Studies on the effect of high temperatures on the thermal stability of various fuels have been continued with emphasis on methods of measuring deposits on tube surfaces. Combustion and electron back scattering are the methods of present choice and an instrument based on the latter principle has been designed. Mathematical models are being devised to represent the various portions of an endothermic fuel system with present emphasis on the development of heat transfer correlations and a model for the dehydrogenation of Decalin. Physical properties for Decalin and JP-8 are included. Calculation of the rate of oxidation of normal octane and HYDROLYNE-M from shock tube studies indicate similar rates of reaction and temperature coefficients.</p>		

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